

GenCore version 5.1.6
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c search, using sw model

il 7, 2004, 17:30:20 ; Search time 4482.16 Seconds
(without alignments)
11294.703 Million cell updates/sec

09-245-198a-1

8

gtgtgagcttggcctgg.....ataaatcatgatttctcttc 1168

NTFY NUC

op 10_0 , Gapext 1.0

0272 seqs, 21671516995 residues

s satisfying chosen parameters: 6940544

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nimum Match 0%

ximum Match 100%

string first 45 summaries

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the number of results predicted by chance to have a

score greater than or equal to the score of the result being prii
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	1168	100.0	1168	6	BD062757	BD062757 A t
2	1168	100.0	1239	10	AF030100	AF030100 M t
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C 4	711	60.9	234182	10	AL603707	AL603707 MC
5	628.6	53.8	1353	6	AX201324	AX201324 Sec
6	628.6	53.8	1353	9	AY358870	AY358870 Hon
7	628.6	53.8	1368	9	AF055872	AF055872 Hon
8	628.6	53.8	1421	6	BD090952	BD090952 Apc
9	624	53.4	1306	9	AF030099	AF030099 Hon
10	614.6	52.6	1373	6	BD062758	BD062758 A t
11	597.8	51.2	1236	6	ARI40407	ARI40407 Sec
12	597.8	51.2	1236	6	BD057124	BD057124 Men
13	566.6	48.5	130254	2	AC136195	AC136195 Rat
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C 18	428	36.6	234801	2	AC118309	AC118309 Rat
19	409.4	35.1	1642	9	BC019047	BC019047 Hon
20	328.4	28.1	1816	9	AY081051	AY081051 Hon
21	304	26.0	218485	2	AC127470	AC127470 Par
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23	278.2	23.8	149555	2	AC126921	AC126921 Bos
24	261	22.3	149736	2	AC126239	AC126239 Fel
25	237.4	20.3	180222	2	AC130192	AC130192 Sus
26	212	18.2	176258	2	AC126925	AC126925 Car
27	111.6	9.6	212093	2	AC126237	AC126237 Car
28	88.4	7.6	7218	6	I66494	I66494 Sequ
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C 39	53.6	4.6	205350	2	AC078946	AC078946 Mus
C 40	53.2	4.6	49430	2	AC100434	AC100434 Mus
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ALIGNMENTS

RESULT 1	BD062757	1168 bp	DNA	linear	PAT 27-
LOCUS	BD062757	A tumor necrosis factor related ligand.			
DEFINITION	BD062757				
ACCESSION	BD062757.1	GI:22608360			
VERSION	JP 2001505407-A/1.				
KEYWORDS	unidentified				
SOURCE	unclassified				
ORGANISM	unclassified				
REFERENCE	1 (bases 1 to 1168)				
AUTHORS	Chicheportriche, Y. and Browning, J. L.				
TITLE	A tumor necrosis factor related ligand				
JOURNAL	Patent: JP 2001505407-A 1 24-APR-2001;				
	BIOMED INC, THE FACULTY OF MEDICINE OF THE UNIVERSITY OF GEN				


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100.0%; Score 1168; DB 10; Length 1239;
ity 100.0%; Pred. No. 1.6e-298;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

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100.0%; Score 1168; DB 10; Length 1239;
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nsensus quality: 22384 bases at least Q20
nated insert size: 210656; sum-of-contents estimation
ality coverage: 0x in Q20 bases; agarose-fp estimation
ality coverage: 7.2x in Q20 bases; sum-of-contents estimation
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: Estimated insert size may differ from sequence length
ee http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
: This is a working draft sequence. It currently
ists of 7 contigs. The true order of the pieces
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rary. Gaps between the contigs are represented as
of N, but the exact sizes of the gaps are unknown.
record will be updated with the finished sequence
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revised.
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8773 118872: gap of unknown length
8873 148924: contig of 30052 bp in length
8925 149024: gap of unknown length
9025 167231: contig of 18207 bp in length
7232 167331: gap of unknown length
7332 189907: contig of 22576 bp in length
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/chromosome="x11"
/clone="RP23-168P5"
60.9%; Score 711; DB 2; Length 203083;
arity 99.6%; Pred. No. 5.5e-177;
onservative 0; Mismatches 0; Indels 3; Gaps 3;
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TGCACTTGTGATGAGGGAAGAGCTGTCTACCTGAAGCTGGACTTGCTGGTGAACGGT 42960
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TGCGCCCTGCCTGCCTGGAAAGATTCTCAGCCACAGCAGCAAGCTCTCCCTGGGGCCC 42900
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TCGGTTGTGTCGACGGTGTCTGGGCTGTGTTCGGCTGTTCGGCCAGGGTCTTCCTTCGG 42840
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[illegible]

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 9 GCTGGCCCTGGGCTCGCTGCAAGAAATTCACAGCCACAGCAGCAAGCTCTCTGGG 541
 10 GCTGGCCCTGGGCTCGCTGGAGGAATTCACAGCCATGCGGGGAGTTCCTCTGGG 612
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 20 AGACTGCCCTCCCTCTAGAGGCTGCTGGGCTCTTTCACGTGTTTCCATCCC 907
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 25 GACCACGTGTTTATTGACTTTGTGCAC----- 968
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34 of the tnfr family useful for treatment and diagnosis of
 35
 36 34.1 GI:22602730

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 14 of the tnf family useful for treatment and diagnosis of
 14
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[illegible]

06:25:09 2004

us-09-245-198a-1.rge

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norvegicus clone RP31-258K6 strain Brown Norway, WORKING
SEQUENCE, 12 ordered pieces.

95.3 GI:31442440
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norvegicus (Norway rat)
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
uses 1 to 130254)
llis A., Ayele K., Beckstrom-Sternberg S.M., Benjamin B.,
ley R.W., Bouffard G.G., Brinkley C., Brooks S., Cariaga K.,
Coleman B., Coleman H., Engle J., Granite S., Guan X.,
J., Haghighi P., Han J., Hansen N., Ho S.-L., Hu P.,
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i.R., Maduro Q.L., Maduro V.B., Margulies E.H., Masiello C.,
i.B., McDowell J., Paquirigan C., Pearson R., Portnoy M.E.,
A., Reddix-Dugue N., Schandler K., Schueler M.G., Shah K.,
C., Stantripop S., Thomas J.W., Thomas P.J., Tsipouri V.,
i.L., Wetherby K.D., Wiggins L., Young A. and Green E.D.
Comparative Sequencing Initiative
uses 1 to 130254)
E.D.
Submission
ted (30-OCT-2002) NIH Intramural Sequencing Center, 8717
ont Circle, Gaithersburg, MD 20877, USA
uses 1 to 130254)
E.D.
Submission

Submission

JOURNAL
COMMENT

Submitted (06-JUN-2003) NIH Intramural Sequencing Center,
Grovmont Circle, Gaithersburg, MD 20877, USA
On Jun 6, 2003 this sequence version replaced gi:27753660.
Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: http://www.nisc.nih.gov
Contact: nisc.zoo@nih.gov
----- Project Information
Center project name: dcf
Center clone name: 258K06

The sequence data in this record represents an 'enhanced'
version of a Phase 2 submission. Specifically, the indicat
order and orientation of each sequence contig has been
established using one or more of the following: read-pair
data from individual subclones, overlaps with neighboring
clones, alignment with available reference sequence (e.g.,
human), and/or confirmation by PCR testing. In addition,
the sequence assembly is based on at least 8x average
coverage in Q20 bases and has been reviewed to rule out
gross misassemblies, the low-quality ends of sequence
contigs have been trimmed away, and each base is associat
with a Phrap-derived quality score.

----- Summary Statistics
Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 128169 bases at least Q40
Consensus quality: 128674 bases at least Q30
Insert size: 150000; agarose-fp
Insert size: 129154; sum-of-contigs
Quality coverage: 10.78x in Q20 bases; agarose-fp
Quality coverage: 12.52x in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 12 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that ha
* provided by the submitter.

* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 10521: contig of 10521 bp in length
* 10522 10621: gap of unknown length
* 10622 13327: contig of 2706 bp in length
* 13328 13427: gap of unknown length
* 13428 28924: contig of 15497 bp in length
* 28925 29024: gap of unknown length
* 29025 39201: contig of 10177 bp in length
* 39202 39301: gap of unknown length
* 39302 41906: contig of 2605 bp in length
* 41907 42006: gap of unknown length
* 42007 70195: contig of 28089 bp in length
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* 77662 94161: gap of unknown length
* 94162 94261: gap of unknown length
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FEATURES
source

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norvegicus clone CH230-154B15, WORKING DRAFT SEQUENCE, 3
ed pieces.

3.8 GI:30521223

GS_PHASE1: HTGS_DRAFT: HTGS_FULLTOP.

norvegicus (Norway rat)

norvegicus

ca; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

ses 1 to 223877)

Marie, Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,
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hausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,
Gibbs, R.A.

Submission

ses 1 to 223877)

ished

K.C.

Submission

ted (06-NOV-2001) Human Genome Sequencing Center, Department
cular and Human Genetics, Baylor College of Medicine, One

REFERENCE AUTHORS TITLE JOURNAL

COMMENT

Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 223877)
Rat Genome Sequencing Consortium.

Submitted (10-MAY-2003) Human Genome Sequencing Center, De

of Molecular and Human Genetics, Baylor College of Medicine,
Baylor Plaza, Houston, TX 77030, USA

On May 10, 2003 this sequence version replaced gi:2508075

The sequence in this assembly is a combination of BAC bases
and whole genome shotgun sequencing reads assembled using
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig
in the feature table below represents a scaffold in the A
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and
by sized gaps filled with Ns to the estimated size. The se
may extend beyond the ends of the clone and there may be s
contigs within a contig-scaffold that consist entirely of
genome shotgun sequence reads. Both end sequences and whole
shotgun sequence only contigs will be indicated in the fea
table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GIOK

Center clone name: CH230-154B15

----- Summary Statistics

Assembly program: Atlas 3.0;

Consensus quality: 214785 bases at least Q40

Consensus quality: 216908 bases at least Q30

Consensus quality: 218593 bases at least Q20

Estimated insert size: 227169; sum-of-contigs estimati

Quality coverage: 7x in Q20 bases; sum-of-contigs esti

* NOTE: Estimated insert size may differ from sequence len

(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_da

* NOTE: This is a 'working draft' sequence. It currently

* consists of 3 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence.

* as soon as it is available and the accession number will

* be preserved.

* 1 221327: contig of 221327 bp in length

* 221328 221427: gap of unknown length

* 221428 222652: contig of 1225 bp in length

* 222653 222752: gap of unknown length

* 222753 223877: contig of 1125 bp in length.

FEATURES

Location/Qualifiers

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/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-154B15"

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/note="wgs contig"

/complement(217607..218056)

/note="clone boundary"

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site:EcoRI

end_sequence:RWBB008TJB"

ORIGIN

Query Match 48.5%; Score 566.6; DB 2; Length 223877;
Best Local Similarity 89.1%; Pred. No. 1.1e-138;
Matches 669; Conservative 0; Mismatches 74; Indels 8; G

Qy 425 CAGGTGCACCTTTCATGAGGAAAGGCTGTCTACTGAAGCTGGACTTCTGGTGAA

Db 147649 CAGGTGCACCTTTCATGAGGAAAGGCTGTCTACTGAAGCTGGACTTCTGGTGAA

GenCore version 5.1.6
copyright (c) 1993 - 2004 Compugen Ltd.

c search, using sw model

il 7, 2004, 17:30:19 ; Search time 2831.52 Seconds
(without alignments)
12318.149 Million cell updates/sec

09-245-198A-1

8
gtgctgagcctggcctgg.....ataaatcatgattctcttc 1168

NTITY_NUC

op 10.0 , Gapext 1.0

13289 seqs, 14931090276 residues

s satisfying chosen parameters: 55026578

th: 0

th: 2000000000

nimum Match 0%

ximum Match 100%

string first 45 summaries

T:*

em_estba:*

em_esthum:*

em_estin:*

em_estmu:*

em_estov:*

em_estpl:*

em_estro:*

em_htc:*

gb_est1:*

gb_est2:*

gb_htc:*

gb_est3:*

gb_est4:*

gb_est5:*

em_estfun:*

em_eston:*

em_gss_hum:*

em_gss_inv:*

em_gss_pln:*

em_gss_vrt:*

em_gss_fun:*

em_gss_nam:*

em_gss_mus:*

em_gss_pro:*

em_gss_rod:*

em_gss_phg:*

em_gss_vrl:*

gb_gss1:*

gb_gss2:*

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
d by analysis of the total score distribution.

SUMMARIES

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.6	918	10	BF577781	602092080	
.8	665	13	BY742288	BY742288	

5	549	47.0	2237	11	AK044387	AK044387 MU
6	533.4	45.7	543	29	CG565104	CG565104 OS
7	519.4	44.5	731	12	BI871711	BI871711 60
8	510	43.7	554	29	CG629394	CG629394 OS
9	507.4	43.4	728	12	BI870393	BI870393 60
10	504	43.2	561	10	AW763237	AW763237 UR
11	497	42.6	533	10	BE628951	BE628951 UU
12	489.6	41.9	650	12	BQ404836	BQ404836 60
13	488.8	41.8	687	13	BQ208433	BQ208433 UI
14	481.2	41.2	584	10	AW917574	AW917574 ES
15	480.8	41.2	939	14	CB849011	CB849011 MR
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17	476	40.8	482	29	CG653257	CG653257 OS
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19	467.8	40.1	948	13	BQ707185	BQ707185 AG
20	464.8	39.8	666	14	CF126539	CF126539 UI
21	456.4	39.1	828	12	BI596681	BI596681 60
22	448.2	38.4	621	29	CG584545	CG584545 OS
23	440.6	37.7	545	14	CB141389	CB141389 X-
24	437.8	37.5	471	9	AA221610	AA221610 my1
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27	422.6	36.2	697	14	CF126932	CF126932 UI
28	421.6	36.1	474	29	CG609156	CG609156 OS
29	421.6	36.1	963	13	BQ671259	BQ671259 AG
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31	419.2	35.9	940	13	BQ884231	BQ884231 AC
32	410.4	35.1	456	13	EX634398	EX634398 BX
33	406.8	34.8	471	29	CG568080	CG568080 OE
34	406.4	34.8	1071	12	BM921213	BM921213 AC
35	400.4	34.3	418	29	CG611020	CG611020 OE
36	393.2	33.7	494	29	CG596702	CG596702 OE
37	390.6	33.4	445	9	AA870722	AA870722 VQ2
38	390.2	33.4	493	29	CG498076	CG498076 OS
39	386.6	33.1	531	29	CG590009	CG590009 OS
40	383.2	32.8	468	29	CG573612	CG573612 OS
41	366.6	31.4	951	13	BQ674188	BQ674188 AG
42	360	30.8	360	10	BE654876	BE654876 UI
43	353.2	30.2	483	29	CG525153	CG525153 OS
44	353	30.2	405	9	AI854476	AI854476 UI-
45	350.6	30.0	414	29	CG599845	CG599845 OS

ALIGNMENTS

RESULT 1
AK020909
LOCUS
DEFINITION
AK020909 1033 bp mRNA linear HTC 20-
Mus musculus adult retina cDNA, RIKEN full-length enriched
clone:A930030D13 product:tumor necrosis factor (ligand)
superfamily, member 12., full insert sequence.

ACCESSION
AK020909.1 GI:12861640
VERSION
HTC; CAP trapper.
KEYWORDS
Mus musculus (house mouse)
SOURCE
Mus musculus
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleo
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murina
1

REFERENCE
AUTHORS
Carninci,P. and Hayashizaki,Y.
TITLE
High-efficiency full-length cDNA cloning
JOURNAL
Meth. Enzymol. 303, 19-44 (1999)
MEDLINE
99279253
PUBMED
10349636
2

REFERENCE
AUTHORS
Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K
Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki
TITLE
Normalization and subtraction of cap-trapper-selected cDNAs
JOURNAL
prepare full-length cDNA libraries for rapid discovery of n
Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE
20499374
PUBMED
11042159

	682	GGCCT	686 -----
Ddb	658	GGCCT	662 -----
RESULT 4	BT742288	665 bp	mRNA linear EST 17
LOCUS	BT742288	Riken full-length enriched,	adult retina Mus musc
DEFINITION	clone AK93003D13 5'	mRNA sequence.	
ACCESSION	BT742288		
VERSION	BT742288.1	GI:27167686	
KEYWORDS	EST.		
SOURCE	Mus musculus	(house mouse)	
ORGANISM	Eukaryota; Metazoa;	Chordata;	Craniata; Vertebrata; Eutele
REFERENCE	Mammalia; Eutheria;	Rodentia;	Sciurognathi; Muridae; Murir
AUTHORS	1 (bases 1 to 665)		
	Kakadoi, I., Furuno, M., Kasukawa, T., Adachi, J.J., Bono, H., Kikuchi, Y., Osato, N., Saito, R., Suzuki, H., Yamanaoka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D.P., Bult, Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsubara, S., Beisel, K.W., Blake, J.A., Bradt, D., Bruscia, V., Chothia, C., Corbani, L.E., Cousins, S., Dalla, E., Draganti, T., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I.J., Jarvis, E.D., Kar Kawai, H., Kawasaki, Y., Kedzierski, R.M., King, B.L., Konaga Kurochin, I.V., Lee, X., Lenhard, B., Lyons, P.A., Maglott, D. Maltais, L., Marchionni, L., McKenzie, L., Mikhi, H., Nagashima Numata, K., Okido, T., Pavan, W.J., Perlea, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandrar Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ring Sandelin, A., Schneider, C., Semple, C.A., Setou, M., Shimada, Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomit Verardo, R., Wagner, L., Wahlesch, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, Z., Wynshaw-Boris, A., Yanagisawa, M., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, Hayatsu, N., Hirozone-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawaji, J., Akizawa, F., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ichihashi, M., Kagawa, I., Miyazaki, A., Sakai, K., Saeki, D., Shii Shinigawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lanc Rogers, J., Birney, E. and Hayashizaki, Y.		
TITLE	Analysis of the mouse transcriptome based on functional ar		
JOURNAL	Of 60,770 full-length cDNAs		
MEBLINE	Nature	420,	563-573 (2002)
PUBMED	22354683		
COMMENT	12466851		
	Contact:	Yoshihide Hayashizaki	
	Laboratory for Genome Exploration Research Group,	RIKEN Ge	
	Sciences Center(GSC), Yokohama Institute		
	The Institute of Physical and Chemical Research (RIKEN)		
	1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa	230-004	
	Tel: 81-45-503-9222		
	Fax: 81-45-503-9216		
	Email: genome-res@sc.riken.go.jp,		
	URL:http://genome.gsc.riken.go.jp/		
	Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Horii, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Koike, M., Kondou, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, K., Nomazaki, R., Ohno, M., Ohsato, N., Saito, R., Saka Sano, H., Saeki, D., Sato, K., Shibata, K., Shiraki, T., Tegar Takeda, Y., Waki, K., Watanahi, A., Muramatsu, M. and Hayashiz Direct Submission		
	Computational Analysis of Full-length Mouse cDNAs Compare Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)		
	Normalization and subtraction of cap-trapper-selected cDNA prepare full-length cDNA libraries for rapid discovery of genes. Genome Res. 10 (10), 1617-1630 (2000).		
	RIKEN integrated sequence analysis (RISA) system--384-for		

ing pipeline with 384 multicapillary sequencer. Genome Res. 1757-1771 (2000)

er-based methods for the mouse full-length cDNA
pedia: real-time sequence clustering for construction of a
ndant cDNA library. Genome Res. 11 (2), 281-289 (2001)

library was prepared and sequenced in Mouse Genome
pedia Project of Genome Exploration Research Group in Riken
Sciences Center and Genome Science Laboratory in RIKEN.
n of Experimental Animal Research in Riken contributed to
mouse tissues.

RNA was provided by Dr. Stefano Gustincich (Department of
ology, Harvard Medical School, 220 Longwood Ave., Boston, MA
USA) whose assistance is gratefully acknowledged.

visit our web site (<http://genome.gsc.riken.go.jp>) for
details.

Location/Qualifiers

1. .665

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="A930030D13"

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/notes="Site 1: Sali; Site 2: BamHI; cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer [5'
GAGAGAGAAGATCCAGAGCTCTTTTCTTTTCTT 3'], cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. cDNA went through one round of normalization
to Rot = 20.0 and subtraction to Rot = 459.0. Second
strand cDNA was prepared with the primer adapter of
sequence [5'GAGAGAGATTCGAGTTAATTAATTAATCCGCCGCCGCC
3']. cDNA was cleaved with XhoI and BamHI. Vector: a
modified pBluescript KS(+) after bulk excision from Lambda
FLC 1. -Retina RNA was provided by Stefano Gustincich,
Department of Neurobiology, Harvard Medical School, 220
Longwood Ave., Boston, MA02115, USA, whose assistance we
gratefully acknowledge."

51.8%; Score 605.6; DB 13; Length 665;
rity 96.4%; Pred. No. 7.7e-147;
nservative 0; Mismatches 22; Indels 2; Gaps 2;

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AGCCATTATGAGGTTCATCTCGGCCAGGACAGATGGAGCAACAGCAGGTGTG 61

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Db 302 CTCCGCTTTGTGCAGGTGTCTGGGCTGTTCGCGCTGCGGCCAGGGTCTTCCCTTCC

QY 608 CGCACCTCCCTGGCTCATCTTAAGGCTGCCCTTCTTAACCTACTTTCGACT

Db 362 CGCACCTCCCTGGCTCATCTTAAGGCTGCCCTTCTTAACCTACTTTCGACT

QY 668 CAAGTTCACCTAGGGGCTTGTCTCTCCAGATTTCCTAAACTTTCCTGGCTCCAG

Db 422 CAAGTTCACCTAGGGGCTTGTCTCTCCAGATTTCCTAAACTTTCCTGGCTCCAG

QY 728 ATCAGACACCTCCCTACCCACCCCTCTCTCCACCCCTCTCTCCACCCCTCTCTGGT

Db 482 ATCAGACACCTTCTACCCCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT

QY 788 TCCTGT-CTCTCTCAAGGCGCAGAGCTTGTTCACATGTTTCCATTTCCACAGAG

Db 542 TCCTGTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT

QY 847 TCCTGTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT

LOCUS AK044387 2237 bp mRNA linear HTC 20-

DEFINITION Mus musculus adult retina cDNA, RIKEN full-length enriched
clone:A930010H24 product:tumor necrosis factor (ligand)
superfamily, member 12., full insert sequence.

ACCESSION AK044387

VERSION AK044387.1 GI:26336423

KEYWORDS HTC; CAP trapper.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele-

AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murin-

TITLE Carninci, P. and Hayashizaki, Y.

JOURNAL High-efficiency full-length cDNA cloning

METH. ENZYMOLOGY Meth. Enzymol. 303, 19-44 (1999)

MEDLINE 99279253

PUBMED 10349636

REFERENCE 2

AUTHORS Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, I.

TITLE Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki:

JOURNAL Normalization and subtraction of cap-trapper-selected cDNA:

MEDLINE Prepare full-length cDNA libraries for rapid discovery of I

PUBMED Genome Res. 10 (10), 1617-1630 (2000)

REFERENCE 3

AUTHORS Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Car-

TITLE Konno, H., Akiyama, J., Nishi, K., Katsunai, T., Tashiro, H., It

JOURNAL Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Hara

MEDLINE Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashi

PUBMED Fujiiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watai

AUTHORS Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., I

TITLE Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki:

JOURNAL RIKEN integrated sequence analysis (RISA) system--384-form

MEDLINE sequencing pipeline with 384 multicapillary sequencer

PUBMED Genome Res. 10 (11), 1757-1771 (2000)

REFERENCE 4

AUTHORS The RIKEN Genome Exploration Research Group Phase II Team

TITLE Functional annotation of a full-length mouse cDNA collecti-

JOURNAL Nature 409, 685-690 (2001)

REFERENCE 5

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QY	1	GGTCTCAGCGCTGGGCGCTGGCGCTGCCTTCGTCCCTCTGCTGTTGGTCTCGTGCT						
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QY	61	GGGAGCTGGGCAACCGCTGTCTGCC---CAGAGAGCTTCTCAGGAGGAGCTGAAC						
Db	477	GGGAGCTGGGCAACCGCTGTCTGCCCAGCAGAGAGCTTCTCAGGAGGAGCTGAAC						
QY	118	GGACCGCGGAGCCCCCTGTAATCCCAGACAGAGAAAGCCACAGGATGTGC						
Db	537	GGACCGCGGAGCCCCCTGTAATCCCAGACAGAGAAAGCCACAGGATGTGC						
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QY	418	GTACTGTCAAGTGCATTTGATCAGGGAAGGCTCTCTACTGAGAGCTGGACTTGC						
Db	837	GTACTGTCAAGTGCATTTGATCAGGGAAGGCTCTCTACTGAGAGCTGGACTTGC						
QY	478	GAA CGGTGTGTCGGCCCTGGCTGCGCTGGAAAGAAATTTCTCAGCCACAGCAGCAAGCT						
Db	897	GAA CGGTGTGTCGGCCCTGGCTGCGCTGGAAAGAAATTTCTCAGCCACAGCAGCAAGCT						
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2237
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Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES
described in Zambrowicz et al (Nature. 1998 Apr 9;392(6674):

06:25:11 2004

us-09-245-198a-1.rst

TTAGGGGCGCTGGTCTCCCGAGTGTGTCCCGAGGTGCCGCTCC 708

94 554 bp DNA linear GSS 02-OCT-2003
515 Mus musculus 129Sv/Ev Mus musculus genomic clone
515, genomic survey sequence.

94
94.1 GI:37453243

sculus (house mouse)

ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
ses 1 to 554)

wicz,B.P., Abuin,A., Ramirez-Solis,R., Richter,L.J.,
t,J., Beltrandi,R.O.,H., Buxton,E.C., Edwards,J., Finch,R.A.,
e,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaing,C.,
W. Jz., Kipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
i,M.J., Van Slightenhorst,I., Vogel,P., Walke,W., Xu,N.,
Person,C. and Sands,A.T.

inase deficiency lowers blood pressure in mice: a gene-trap
to identify potential targets for therapeutic intervention
Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)

it: Zambrowicz Bp

unk

in Genetics Incorporated

Research Forest Drive, The Woodlands, TX 77381, USA

materials@lexgen.com

rap sequence tag generated by 3' RACE from mouse ES cells as
bed in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)

Gene Trap.

Location/Qualifiers

1. .554

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43.7%; Score 510; DB 29; Length 554;

arity 98.9%; Pred. No. 6.8e-122;

conservative 0; Mismatches 3; Indels 3; Gaps 3;

GTGGATGGGACAGTGAGTGGCTGGGAAGAGACCAAAATCAACAGCTCCAGCCCTCT 360

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TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 420

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TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 480

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 185

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 539

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 245

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 599

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 305

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 659

TTACGACCCGACATTTGGGGAATTTACAGTCATCAGGCTGGGCTTACTACTGTA 365

QY 660 GACTCTTTCAAGTTCACTGAGGGGCTTGTCTCTCCAGATTCTTAAACTTTCCCT
Db 366 GACTCTTTCAAGTTCACTGAGGGGCTTGTCTCTCCAGATTCTTAAACTTTCCCT

QY 720 CCAGGAGATCACCACACCTCCCTACCCACCCACCCACCTCTCCACCCCTCGCTG
Db 426 CCAGGAGATCACCACACCTCCCTACCCACCCACCCACCTCTCCACCCCTCGCTG

QY 780 TGGTCCAGTCTCTGT-CTCTCTCAAAGGAGCCAGAGCTTGTTCACATG-TTTCG
Db 486 TGGTCCAGTCTCTGTCTCTCAAAGGAGCCAGAGCTTGTTCACATGTTTTCG

QY 838 ACAGACGTA 846

Db 546 ACAGACGTA 554

RESULT 9

BI870393

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

COMMENT

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AP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 al Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Gene Index
 ished (1997)
 t: Robert Strausberg, Ph.D.
 cgapbs-r@mail.nih.gov
 lone is available royalty-free through LLNL; contact the
 Consortium (info@image.llnl.gov) for further information.
 83048
 imer: -40RP from Gibco
 uality sequence stop: 437.
 Location/Qualifiers
 1. .533
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 polylinker; 1st strand cDNA was prepared from mammary
 gland tissue from a lactating female, and was then primed
 with a Not I - oligo(dT) primer. Double-stranded cDNA was
 ligated to Eco RI adaptors (Pharmacia), digested with Not
 I and cloned into the Not I and Eco RI sites of the
 modified pTV73 vector. Library is normalized. Library
 was constructed by Bento Soares and M. Fatima Bonaldo."

42.6%; Score 497; DB 10; Length 533;

arity 99.4%; Pred. No. 1.7e-118;

conservative 0; Mismatches 0; Indels 3; Gaps 3;

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 |||||
 GTGCTGGCCCTCCCTGCTGGAGAAATTCACGCCAGCAGCAAGCTCTCTGG 60
 |||||
 CAGCTCCGTTTGTGCCAGTGTCTGGGCTGTGGCGCTGGCGCCAGGCTCTTCCT 600
 |||||
 CAGCTCCGTTTGTGCCAGTGTCTGGGCTGTGGCGCTGGCGCCAGGCTCTTCCT 120
 |||||
 ATCCGACCCCTCCCTGCTGCTAAAGCTGCTGCTGCTGCTGCTGCTGCTGCT 660
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 ATCCGACCCCTCCCTGCTGCTAAAGCTGCTGCTGCTGCTGCTGCTGCTGCT 180
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 TTTCAGGTTCACTGAGGGGCTTGTCTCCAGATTCCTTAACTTTTCCCTGGCTC 720
 |||||
 TTTCAGGTTCACTGAGGGGCTTGTCTCCAGATTCCTTAACTTTTCCCTGGCTC 240
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 AGCATCACACACCTCCCTACCCACCCCACTCTCCACCCCTGCTGCTGCTGCT 780
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 AGCATCACACACCTCCCTACCCACCCCACTCTCCACCCCTGCTGCTGCTGCT 300
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 CAGTCCCTGT-CTCTCTCAAGGAGCAGAGCTGTTTCAATG-TTTCATTTCCA 838
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 CAGTCCCTGTCTCTCTCAAGGAGCAGAGCTGTTTCAATGTTTTCATTTCCA 360
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 CGTATCCTTGTCTTTC-TTACATCCCATCCACAGACTATCCACTCACTAGC 897
 |||||
 CGTATCCTTGTCTTTC-TTACATCCCATCCACAGACTATCCACTCACTAGC 420
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 |||||
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 TGTGACACGAGCATTGAGATGGGCTGGAGCTGGTGGAGGAGCCAGAG 533
 |||||

BG404836
 LOCUS
 DEFINITION 60242016QF1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:4527C
 mRNA sequence.
 BG404836
 BG404836.1 GI:13298284
 EST.
 Mus musculus (house mouse)
 Mus musculus
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murin
 NIH-MGC <http://mgi.nci.nih.gov/>.
 1 (bases 1 to 650)
 National Institutes of Health, Mammalian Gene Collection (
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: The Cepko Laboratory
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information ca
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: LHAM10435 row: h column: 15
 High quality sequence stop: 468.
 Location/Qualifiers
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 /clone_lib="NIH_MGC_94"
 /note="Organ: eye; Vector: pCMV-SPORT6; Site 1: 1
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 Average insert size 3.3 Kb. Library enriched for
 full-length clones and constructed by Life Techn
 Note: this is a NIH_MGC Library."

ORIGIN

Query Match 41.9%; Score 489.5; DB 12; Length 650;
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 QY 351 CCAGCCCTCTCGGTACGACCGCCAGATTGGGGAATTTACAGTCATCAGGGCTGG
 DB 85 CCAGCCCTCTCGGTACGACCGCCAGATTGGGGAATTTACAGTCATCAGGGCTGG
 QY 411 ACTACCTGTACTGTTCAGTGCACCTTTGATGAGGGAAGGCTGTCTACCTGAAGCT
 DB 145 ACTACCTGTACTGTTCAGTGCACCTTTGATGAGGGAAGGCTGTCTACCTGAAGCT
 QY 471 TGCTGGTGAACGGTGTCTGGCCCTCGCTGGCTGGCTGGGAAGAAATTTCTCAGCCACAGC
 DB 205 TGCTGGTGAACGGTGTCTGGCCCTCGCTGGCTGGCTGGGAAGAAATTTCTCAGCCACAGC
 QY 531 GCTCTCCCTGGGCCCGAGCTCCGTTTGTGCGCAGGTGTCTGGGCTGTGGCGCTGGC
 DB 265 GCTCTCCCTGGGCCCGAGCTCCGTTTGTGCGCAGGTGTCTGGGCTGTGGCGCTGGC
 QY 591 GGTCTTCCTTCGATCCGACCCCTCCCTGGGCTCATCTTAAAGCTGCCCCCTTC
 DB 324 GGTCTTCCTTCGATCCGACCCCTCCCTGGGCTCATCTTAAAGCTGCCCCCTTC
 QY 651 CCTACTTTGAGTCTTTTCAAGTTTCACTGAGGGGCTTGTCTCTCCAGATTCTTTA
 DB 384 CCTACTTTGAGTCTTTTCAAGTTTCACTGAGGGGCTTGTCTCTCCAGATTCTTTA

ses 1 to 584)
H., Glodek, A., Chandra, I., Mason, T.M., Quackenbush, J.,
age, A.R. and Adams, M.D.
name Project: Generation of a Rat EST (REBT) Catalog & Rat
index
ished (1998)
t: Lee, NH
stitute for Genomic Research
Medical Center Drive, Rockville, MD 20850, USA
301)-838-3529
301)-838-0208
nhlee@tigr.org
lone is available through the ATCC, contact the ATCC
3-365-2700 for further information
imer: M13 Reverse.
Location/Qualifiers
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41.2%; Score 481.2; DB 10; Length 584;
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onservative 0; Mismatches 46; Indels 5; Gaps 2;
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CCAAATCAACAGCTCCAGCCCTCTGGCTACGACCGCCAGATTGGGAATTACA 391
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TCAGGGCTGGGCTCTACTACCTGTACTGTCTAGGTGCACTTTGATGAGGAAAGGCT 451
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CAGCCACAGCAGCAGCTCTCTGGGCCCCAGCTCCGTTTGTGCGAGGTGTCTGGG 301
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CTGCCCCCTTCTTACCTACTTTGGACCTTTCAAGTTCACTGAGGGCTTGTCTC 421
TAGATTCTTTAACTTTCCCTCTGGCTCCAGGACATCACACACCTCCCTATCCC 747
TAGTTCTTTAACTTTAACTTTAACTTTAACTTTAACTTTAACTTTAACTTTAACTTT 481
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TCCACTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 541
TAGAGCTTGTTCATATG-TTTTCCATTTCCAGACGATATCC 849

542 AGCCAGNGCTTGATGACATGTTTTCATTCCACAGACATATTC 584
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LOCUS
DEFINITION
MRA-0673 MOUSE ADULT RETINA Mus musculus cDNA 5', mRNA seq
ACCESSION
CB849011
VERSION
CB849011.1 GI:34379595
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murin
1 (bases 1 to 939)
Yu, J., Farjo, R., MacNee, S.P., Baehr, W., Stambolian, D.E. and
Swaroop, A.
Annotion and analysis of 10,000 expressed sequence tags
developing mouse eye and adult retina
Genome Biol. 4 (10), R65 (2003)
2281944
14519200
PUBMED
COMMENT
Contact: Swaroop, A.
Department of Ophthalmology and Visual Sciences
Kellogg Eye Center, University of Michigan
540 KEC, 1000 Wall St., Ann Arbor, MI 48105, USA
Tel: 734 615 2246
Fax: 734 647 0228
Email: swaroop@umich.edu.
FEATURES
Location/Qualifiers
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Query Match 41.2%; Score 480.8; DB 14; Length 939;
Beat Local Similarity 90.4%; Pred. No. 3.7e-114;
Matches 553; Conservative 0; Mismatches 52; Indels 7;
QY 225 GGAAGGCGCGCTCTGGCGAGCTATTCAGCCCATTTATGAGGTTCATCTCGGCCF
DB 25 GGAAGGCTGTCTCGCTCAGGTACCGCTCCGGAATTCGCCGGTCCGACCCAGCGF
QY 285 AGGATGGAGCACAAGCAGGTGTGGATGGACAGTGAGTGGCTGGGAGAGACCAAF
DB 85 AGGATGGAGCACAAGCAGGTGTGGATGGACAGTGAGTGGCTGGGAGAGACCAAF
QY 345 ACAGCTCCAGCCCTCTGCGCTACGACCGCCAGATTGGGGAATTTACAGTCATCAG
DB 145 ACAGCTCCAGCCCTCTGCGCTACGACCGCCAGATTGGGGAATTTACAGTCATCAG
QY 405 GGCTCTACTACTGTACTGTCTGAGGTGCACTTTGATGAGGGAAGGCTCTTACCTC
DB 205 GGCTCTACTACTGTACTGTCTGAGGTGCACTTTGATGAGGGAAGGCTCTTACCTC
QY 465 TGGACTTGTCTGCTGAAAGGCTGTCTGGCCCTGCGCTGCTGCGGAATTTCTCAGCC
DB 265 TGGACTTGTCTGCTGAAAGGCTGTCTGGCCCTGCGCTGCTGCGGAATTTCTCAGCC
QY 525 CAGCAAGCTCTCTGGGCCCCAGCTCCGTTTGTGCGGAGGTGTCTGGGCTGTTCGCC
DB 325 CAGCAAGCTCTCTGGGCCCCAGCTCCGTTTGTGCGGAGGTGTCTGGGCTGTTCGCC
QY 585 GGCAGGAGGTCTTCCCTTCGGATTCGCGACCCCTCCCTGGGCTCATCTTAAGGCTGC
DB 385 GGCAGGAGGTCTTCCCTTCGGATTCGCGACCCCTCCCTGGGCTCATCTTAAGGCTGC
QY 645 TCCTAACCTACTTTGGATCTCTTTCAAGTTCACTGAGGGGCTTGTCTCTCCAGATTT

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TTNCTGGCTCCAAAGGAGCATCACACACCTGCTTACCCACCCCTCTCTCCA 564
T-CGCTGCTCCTTGGTCCAGTCC--TGTCTCTCTCTCAAGGCA--GCCAGAGCT 817
TNNCTGCTCCTTGGGACCANTCCTGTTCTCTCTCTNCTNAAAGGCANNNCAANAGCT 624
ACATGTT 829
ACATGTT 636

Pril 7, 2004, 23:15:14
secs

CTAACCTACTTTGGACTCTTTCAGTTCACCTGAGGGCCCTTGCTCTCCAGATTCCT 660

bladder and central nervous system cancer, melanoma, breast, ovarian, renal, colorectal, uterine, prostate, and thyroid cancer.

our polypeptides

BP; 257 A; 443 C; 389 G; 264 T; 0 U; 0 Other;

53.8%;	Score 628.6;	DB 3;	Length 1353;
ity 76.2%;	Pred. No. 1.2e-159;		
nservative	0;	Mismatches 219;	Indels 73; Gaps 10;

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CCGGGCATCGCTGTCGCCCGCAGAGCGCTGCCAGGAGAGCTGGTGCAGAGAG 249
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 .TATTGCAGCCCATTTATGAGGTTTCATCTCTGGCCAGACAGAGATGGAGCAACAACA 301
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CGACCGCCAGATTGGGGAATTACAGTCATCAGGCTGGGCTCTACTACCTGTAC 421

GGTGCACTTTGATGAGGGAAAGCTGTCTACCTGAAGCTGGACTTGCTGGTGAAC 481

'GCTGGCCCTGCGTGCCTGGGAAGAAATTCACAGCCACAGCAGCAAGCTCTCCTGGG 541

601 GCTCCGTTTGTGCCAGGTGTCTGGGCTGTGCCGCTGGCGCCAGGGTCTTCCCTT

AGCTCCGCCCTCTGCCAGGATCTGGGCTGTTGGGCTCTGCGGCCAGGGTCTCTCCCTCCTG 729

TCGACACCCCTCCCCCTGGGCCCCCATCTCAAGGCGTGGCCCCCTTCCCTCACCCTACTTCGGA 789

TCAGGTTCACTGAGGGGCCCTTGCTCTCCAGATTCCTTAAACTTTCCTGGCTCC 721

CCAGGTTCACTAGGSGCCCTGGTCTCCCGGCAGTGGTCCACAGGCTGCCGGGTCC 849
;CATCACACACCTCCCTATCCCCACCCCACTCCTCCACCCCCCTC-GCTGCTCCTT 780

-CTCGACAGCTCTCTGGCACCCGGTCCCTCTGCCCCACCCCTCAGCCGCTCTTT 904
 AGTCCTGTCTCTCC--TCAAAGGCAGCCAGAGCTTGTTCACATGTTTCCATTCC- 837

2AGACCTGCCCCCTCCCTCTAGAGGCTGCCTGGGCCCTGTTACGTTGTTTTCCATCCC 964
 --ACAGACGTAACCTTGCTCTTCTTAACATCCCATCCACCAAACTATCCACCTC 891

AAATACAGTATTCCCACTTATCTTACAACACTCCCCCAGCGCCCACTCTCCACCTC 1024
3TCCCAAAGCCCTAC-----TTATCCCTGACTCCCCCACT 936

3CTCCCCAATCCCTGACCCCTTGAGGCCCCAGTGATCTGACTCCCCCTGGCCA 1084
2GACCACGTGTTTATTGACTTTGTGCAC----- 968

1085	Db	CAGACCCCCCAGGTCATGTGTTCATCTGTACTCTGTGGCAGAGGATGGGTCCAGAG
969	QY	-----CAGGCACTCAGATGGGTGGACCTGGTGGCAGGAAGCCAGAGAACTCTGGGA
1145	Db	CAC TTCAGGCACTAAGAGGGGGCTGGACCTGGCGGCAGGAAGCCAAAGAGACTGGGGC
1024	QY	GCACAGAGTTCCCACTGTGAGGGGGAAGAGCTGGGGAACAGCTCCCTCCCTGGA
1205	Db	GCCAGGAGTTCCCAATGTGAGGGGCGAGA--AACAGACAAGCTCTCCCTTTCAGAG
1080	QY	CCTGTGATTTTCAAA--AGATACTATTTTTATTATTATTGTGCACAAAATGT---TT
1264	Db	CCTGTGATTTTTAAACAGATATTATTTTATTATTATTGTGCACAAAATGTTGAT
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RESULT 5
ABK40255
ID ABK40255 standard: cDNA: 1353 BP.

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AC	
XX	15-JUL-2002 (first entry)
DT	

XX
DE
XX
XW

KW leukaemia; neuronal disorder; stromal disorder; blastocoelec diso.
 KW inflammatory disorder; immune disorder; angiogenic disorder;
 KW gene therapy; cytostatic; neuroprotective; gene; ss.

OS Homo sapiens.

PN WO200153486-A1.

PD 26-JUL-2001.

11-FEB-2000: 2000WO-US003565.

XX
08-MAR-1999.
99W0-IIS005028

PR 11-MAR-1999; 99US-0123972P.
PR 11 MAY 1999; 99US 0123450P

PR 02-JUN-1999; 99WO-US012252.

PR 22-JUN-1999; 99US-0140653P.

PR 26-JUL-1999; 99US-0145698P.

FR 28-JUN-1999; 99US-0149395P;
PR 17-AUG-1999; 99US-0149395P;

PR 31-AUG-1999; 99US-0151689P.
PR 01-SEP-1999: 99WO-US020111.

PR 15-SEP-1999; 99WO-US021090.
PR 30-NOV-1999; 99WO-US028313

PR 01-DEC-1999; 99WO-US028301.
PR 01-DEC-1999; 99WO-US028301.
PR 01-DEC-1999; 99WO-US028301.

PR 05-JAN-2000; 2000WO-US000219.

PA (GETH) GENENTECH INC.

PI Ashkenazi AJ, Goddard A, Godowski PJ, Gurney AL, Hillan KJ;

PI Marsters SA, Pan J, Pictt RM, Roy MA, Smith V, Stone DM;
PI Watanabe CK. Wood WI:

XX
WBT - 2002-2003 JAM 36/295502

DR P-PSDB; AAU86129.

PT Thirty five nucleic acids encoding PRO polypeptides, useful for t
PT benign or malignant tumors, leukemias and lymphoid malignancies,
PT inflammatory, angiogenic and immunologic disorders.

vention relates to the isolation of novel human PRO and the polynucleotide sequences encoding them. The PRO agonists, antagonists or anti-PRO antibodies are useful for gn or malignant tumors (e.g. renal, kidney, bladder, leukemias and lymphoid malignancies, other disorders such as glial, astrocytal, hypothalamic, glandular, macrophagal, glioma, osteocytic disorders, inflammatory, immune and angiogenic disorders). The polynucleotide sequences are also useful in gene therapy. The polynucleotide encode for the human PRO polypeptides of the invention

BP: 257 A; 443 C; 389 G; 264 T; 0 U; 0 Other;

53.8%; Score 628.6; DB 6; Length 1353;
arity 76.2%; Pred.No.1.2e-159;
onservative 0; Mismatches 173; Gaps 10;

TGTGACCTGGGCCTGGCGTCCGCCCTGCTGGCCTCTGTCGTGCTGGTGTCAGCCTG 61
TTGGACCCTGGGCCTGGCGTCCGCCCTCTGTCGCCGTGGTCACTTTG 189
TAGAGCTGGGAACCGCTGTCTGCCAGGAGCCCTTCTCAGAGGAGCTGCAGCAGAGGAC 121
AGCCGGGCATCGCTCTCGCCCCAGGAGCCCTGCCAGGAGAGCTGGTGGCAGAGGAG 249
GGGAGCCCCCTTGAACTGAATCCCAGA CAGAGGAAGCACGAGATGTGTACTCTTC 181
TAGAACCCGCTCGA ACTGAATCCCAGACAGAAAAGCCAGATCTCGGCCCTTC 309
TAACAACACTAGTCGGCCTCGMAAGTGTCTATAAGSCCGAAGCGCGCCTCGC 241
TAACCAACTAGTTGGCCTCGAAGTGTACCTTAAGGCCGAAAAACAGGCTCGA 369
GCTATTTCAGCCCATPATGAGGTTTCATCTCGGCCAGSACAGGATGGAGCACAGCA 301
GCGATCGACGCCATTATGAAGTTTCATCCACGACCTGGACAGSACGAGCGCAGGCA 429
TGTGGATGGGACAGTAGTGGCTGGGAAGAGACCAAATCAACAGCTCCAGCCCTCTG 361
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--- -- CCTGACAGCTCTCTGGGAC ACCGGTCCCTCTGCCCCACCTCAGCGCTCTTT 904
CCAGTCTGTCTCTCC--TCAAAGGACCGCAGAGCTTTGTTCACATGTTTCATTCC- 837

905	GCTCCGAGACCTGCCCTCCCTCTAGAGGCTGCTGGGCTCTGTTCAAGTGTCTTC	Dd
838	-----ACAGACGTATCTCTTCTTTAAATCCCATCCACACCAACTATCC	Qy
965	ACATAAATACAGTATTCCTTAACTCTCCCTCCACCGCCCACTCTCC	Dd
892	ACTAGTCTCCCAAGGCCCTTAC-----TTATCCTGACTCCCCCACC	Qy
1025	ACTAGCTCCCAATCCCTGACCCCTTTGAGGCCCCAGTGATCTCGACTCCGCCCT	Dd
937	CACCGCACCACTGTTTATTATGACTTTGTGCAC-----	Qy
1085	CAGACCCCCAGGTCATTGTGTTCACTGTACTCTGTGGGCAAGATGGGTCCAGAA	Dd
969	-----CAGGCACTGAGATGGGCTGGACCTGGTGCGAGGAAGCCAGAGAACTCTGGG	Qy
1145	CAC TTCAGGCAC TAAGAGGGGCTGGACCTGGCGGCAGGAAGCAAGACACTGGG	Dd
1074	GCCAGAGTTCCTCCAACTGTCAGGGGGGAGAGACTGGGGACAAGCTCCTCCCTCGA-	Qy
1205	GCCAGGAGTTCCTCCAAATGTGAGGGGGGAGA-AACAAGACAAGCTCTCCTCTTGAG	Dd
1080	CTGTGGATTTTGAAA--AGATACATTTTTTATTATTATTGTGACAAAATGT---	Qy
1264	CTGTGGATTTTAAACAGATATTATTTTATTATTATTGTGACAAAATGTGTA	Dd
1135	GGATATTAAAGAGAATAAATCATGA 1159	Qy
1324	GGATATTAAATAGAAATAGTCATAA 1348	Dd

RESULT 6
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ID AAX56000 standard: DNA: 1421 BP.

XX
AC AAX56000:

AA	15-JUL-1999	(first entry)
DT		

Human tumour necrosis factor Apo-3 ligand polynucleotide sequence
XX
DE
XX
Human; tumour necrosis factor; Apo-3 ligand; lymphotoxin; apopto
KW
KW
NF-kappaB-dependent transcription; JNK/SAPK-dependent response;
KW
KW
ss.

XX Homo sapiens.

XX	Key	Location/Qualifiers
FH	CDS	92. .841
FT		

```
ET / *tag= a
ET /product= "Apo-3 liqand"
```

XX PN WO9919490-A1.

XX
PD
22-APR-1999.

09-OCT-1998:

AA
PR 10-OCT-1997; 97US-0062037P.

FK XX
10-DEC-1968, 2703-000000Z

XX
PA (GEIH) GENENTECH INC.

XX
 FBI MEMPHIS
 ASSASSINAT RO, MATSUCI OSA,
 44-1987-1000

DK WFI; 1935-287502/23.
DR P-PSDB: AAY09369.

XX New human ApoB-100 (a tum

PS Claim 18: Fig 1: 74pp; English

CC The present sequence encodes

Conservative 0; Mismatches 220; Indels 73; Gaps 10;

1024 GCCAGAGTTCCCACTGTGAGGGGGAAGAGCTGGGGAACAAGCTCCTCCCTGGA--
 1165 GCCAGGAGTTCCCAATGTGAGGGGCGAGA-AACAAGACAAGCTCCTCCCTTGAG/
 1080 CCTGTGGATTTTGAAA--AGATACATATTTTATTATTATTGACAAAATCT---
 1224 CCTGTGGATTTTAAACAGATATTTATTATTATTATTATTGACAAAATGTTGA/
 1135 GGATATTAAAGAGAAATAATCA 1156
 1284 GGATATTAAATAGATTAAGTCA 1305

RESULT 8
 ACC57901
 ID ACC57901 standard; cDNA; 1306 BP.
 XX ACC57901;
 AC ACC57901;
 DT 11-AUG-2003 (first entry)
 XX Human TWEAK coding sequence.
 DE Human; TWEAK; tumour necrosis factor; ligand; cytostatic;
 KW immunomodulator; osteopathic; gene; ss.
 XX Homo sapiens.
 XX Key Location/Qualifiers
 FT CDS 18..767
 FT /*tag= a
 FT /product= "Human TWEAK"
 XX WO2003040307-A2.
 PN 15-MAY-2003.
 XX 25-JUL-2002; 2002WO-US023782.
 XX 27-JUL-2001; 2001US-0307838P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX Hilbert DH, Rosen CA;
 PI WPI; 2003-430659/40.
 DR P-PSDB; ABR42315.
 XX New heteromultimeric complex having a first polypeptide member o
 PT tumor necrosis factor (TNF) ligand family, and a second differen
 PT of TNF ligand family, useful for treating cancer, osteoporosis o
 PT autoimmune disease.
 XX Disclosure; Page 367-368; 388pp; English.
 XX The present sequence is that of a polynucleotide encoding human
 CC The invention relates to compositions comprising heterotrimeric
 CC of tumour necrosis factor (TNF) ligand family members, and their
 CC the detection, prevention and treatment of disease. In one embod
 CC the heterotrimeric complex comprises full-length or extracellular
 CC portions of TWEAK and full-length or extracellular portions of o
 CC ligand family members, preferably VEGF or VEGF-SV. The heterotri
 CC complexes of the invention are useful for treating an autoimmune
 CC cancer or osteoporosis, and particularly for inhibiting cancer c
 CC proliferation, increasing B cell proliferation, or inducing apop
 CC T cells

Query Match 53.4%; Score 624; DB 7; Length 1306;
 Best Local Similarity 76.0%; Pred. No. 2.1e-158;
 Matches 929; Conservative 0; Mismatches 220; Indels 73;

06:25:10 2004

Conservative 0; Mismatches 220; Indels 73; Gaps 10;

1024 GCCAGAGTTCCCACTGTGAGGGGGAAGAGCTGGGGAACAAGCTCCTCCCTGGA--
 1165 GCCAGGAGTTCCCAATGTGAGGGGCGAGA-AACAAGACAAGCTCCTCCCTTGAG/
 1080 CCTGTGGATTTTGAAA--AGATACATATTTTATTATTATTGACAAAATCT---
 1224 CCTGTGGATTTTAAACAGATATTTATTATTATTATTATTGACAAAATGTTGA/
 1135 GGATATTAAAGAGAAATAATCA 1156
 1284 GGATATTAAATAGATTAAGTCA 1305

RESULT 8
 ACC57901
 ID ACC57901 standard; cDNA; 1306 BP.
 XX ACC57901;
 AC ACC57901;
 DT 11-AUG-2003 (first entry)
 XX Human TWEAK coding sequence.
 DE Human; TWEAK; tumour necrosis factor; ligand; cytostatic;
 KW immunomodulator; osteopathic; gene; ss.
 XX Homo sapiens.
 XX Key Location/Qualifiers
 FT CDS 18..767
 FT /*tag= a
 FT /product= "Human TWEAK"
 XX WO2003040307-A2.
 PN 15-MAY-2003.
 XX 25-JUL-2002; 2002WO-US023782.
 XX 27-JUL-2001; 2001US-0307838P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX Hilbert DH, Rosen CA;
 PI WPI; 2003-430659/40.
 DR P-PSDB; ABR42315.
 XX New heteromultimeric complex having a first polypeptide member o
 PT tumor necrosis factor (TNF) ligand family, and a second differen
 PT of TNF ligand family, useful for treating cancer, osteoporosis o
 PT autoimmune disease.
 XX Disclosure; Page 367-368; 388pp; English.
 XX The present sequence is that of a polynucleotide encoding human
 CC The invention relates to compositions comprising heterotrimeric
 CC of tumour necrosis factor (TNF) ligand family members, and their
 CC the detection, prevention and treatment of disease. In one embod
 CC the heterotrimeric complex comprises full-length or extracellular
 CC portions of TWEAK and full-length or extracellular portions of o
 CC ligand family members, preferably VEGF or VEGF-SV. The heterotri
 CC complexes of the invention are useful for treating an autoimmune
 CC cancer or osteoporosis, and particularly for inhibiting cancer c
 CC proliferation, increasing B cell proliferation, or inducing apop
 CC T cells

Query Match 53.4%; Score 624; DB 7; Length 1306;
 Best Local Similarity 76.0%; Pred. No. 2.1e-158;
 Matches 929; Conservative 0; Mismatches 220; Indels 73;

Qy	1024	GCAGAGTGTCCCACTGTGAGGGGGAAGAGCTGGGGAACAAGCTCTCTCCCTGCA--	
Db	1165	GCAGAGTGTCCCAAAATGTGAGGGGGGAGA-AACAAGACAAGCTCTCTCCCTTGAGA-	
Qy	1080	CCTGTGGATTTTCAAA--AGATACTATTTTATTTATTTATTTGTGACAAAATGT---T	
Db	1224	CCTGTGGATTTTAAACACAGATATTTTATTTATTTATTTATTTGTGACAAAATGTTGAT	
Qy	1135	GGATATTAAAGAGAATAAATCA 1156	
Db	1284	GGATATTAAAGATAAAGTCA 1305	
RESULT 9			
ADC35205			
ID	ADC35205	standard; cDNA; 1306 BP.	
XX	ADC35205;		
AC	ADC35205;		
XX	18-DEC-2003 (first entry)		
DT	18-DEC-2003 (first entry)		
XX	Human cDNA encoding TNF ligand family member #12.		
DE	ss; gene; human; tumour necrosis factor; TNF ligand; endokine alp		
KW	excessive bone resorption disorder; osteoporosis; Paget's disease		
KW	arterial calcification.		
OS	Homo sapiens.		
XX	US2003100074-A1.		
PN	29-MAY-2003.		
XX	15-AUG-2002; 2002US-00218547.		
PF	16-AUG-2001; 2001US-0312542P.		
PR	30-OCT-2001; 2001US-0330761P.		
XX	(YUGG/) YU G.		
PA	(NIJ/) NI J.		
PA	(ROSE/) ROSEN C A.		
PA	(NARD/) NARDELL B.		
PI	Yu G, Ni J, Rosen CA, Nardelli B;		
XX	WPI; 2003-696072/66.		
DR	P-PSDB; ADC35206.		
XX	New Endokine alpha gene useful for preparing a composition for tr		
PT	disease associated with excessive or insufficient bone resorption		
PT	osteoporosis, Paget's disease or arterial calcification.		
XX	Disclosure; SEQ ID NO 23; 145pp; English.		
PS	The invention relates to an isolated nucleic acid molecule encodi		
XX	tumour necrosis factor family ligand. A composition comprising th		
CC	isolated antibody or its fragment is used for treating an individ		
CC	need of decreased level of endokine alpha activity. The endokine		
CC	polypeptide present in a heterotrimeric complex is used for treat		
CC	individual having a disorder associated with excessive bone resor		
CC	e.g. osteoporosis, Paget's disease or arterial calcification. Tre		
CC	individual having a disorder associated with insufficient bone re		
CC	comprises administering an endokine alpha antagonist, which is th		
CC	antibody that binds specifically to endokine alpha polypeptide. T		
CC	present sequence represents a cDNA encoding a tumour necrosis fac		
XX	family ligand.		
XX	Sequence 1306 BP; 247 A; 434 C; 368 G; 257 T; 0 U; 0 Other;		
SQ			
Query Match	53.4%;	Score 624;	DB 9; Length 1306;
Best Local Similarity	76.0%;	Pred. No. 2.le-158;	
Matches 929;	Conservative 0;	Mismatches 220;	Indels 73;

QY	969	-----CAGGCACTCAGATGGCTGGACCTGGTGGCAGGAAGCCAGAGACCTGGGG	
Db	1088	CACCTTCAGGCACTAAGAGGGGGCTGGACCTGGCGGCAAGGCCAAGAGACCTGGGG	
QY	1024	GCCAGAGCTCCCAACTGTGAGGGGGAAGAGCTGGGGCAAAAGCTCTCCCTCGGA--	
Db	1148	GCCAGAGCTCCCAACTGTGAGGGGGAAGAGCTGGGGCAAAAGCTCTCCCTCGGA--	
QY	1080	CTGTGGATTTTGAAGAATCTATTTT 1108	
Db	1207	CTGTGGATTTTGAAGAATCTATTTT 1235	
RESULT 14			
AAAX23424			
ID	AAAX23424	standard; DNA; 1030 BP.	
AC	AAAX23424;		
XX			
XX	18-JUN-1999	(first entry)	
DE	Human TNRL3 DNA.		
XX			
KW	Tumour necrosis factor receptor; signal transducer molecule; TNF		
KW	developmental abnormality; gestational abnormality; prostate c		
KW	AP06; AP08; AP09; TNRL-1; TNRL-3; diagnosis; treatment; therapy; c		
KW	cytoplasmic domain; immunogen; antibody preparation; breast carc		
KW	apoptosis; human; ss.		
OS	Homo sapiens.		
XX			
Key	Location/Qualifiers		
PH	1..627		
FT	/*tag= a		
FT	/product= "TNRL3"		
FT			
XX	WO9911791-A2.		
PN			
XX			
PD	11-MAR-1999.		
XX			
PF	04-SEP-1998; 98WO-US018393.		
XX			
XX	05-SEP-1997; 97US-00924634.		
PR			
XX	(UNIW) UNIV WASHINGTON.		
PA			
XX			
PI	Chaudhary PM;		
XX			
XX	WPI; 1999-205191/17.		
DR	P-PSDB; AAW93590.		
DR			
PT	New Tumor Necrosis Factor family receptor polypeptides and ligand		
PT	useful for diagnosis and treatment of prostate cancer and develop		
PT	or gestational abnormalities.		
XX			
PS	Example VII; Fig 13A; 156pp; English.		
XX			
CC	This invention describes isolated Tumor Necrosis Factor (TNF) fa		
CC	receptor polypeptides: APO4, APO6, APO8 and APO9 or their active		
CC	fragments, and isolated TNF related ligands 1 and 3 (TNRL1 and 1		
CC	their active fragments. APO4 is useful for diagnosing prostate c		
CC	determining levels of APO4 in an individual. Prostate cancer can		
CC	treated using APO4 selective binding agents linked to a therap		
CC	moieties. APO4 polypeptides are also useful for identifying select		
CC	binding agents, useful in diagnosis/treatment of disease by bindi		
CC	agents to the polypeptide/active fragment which is extracellular		
CC	expressed on the cell surface. The binding is preferably perform		
CC	vivo. APO4 polypeptides/ active fragments are also useful for sc		
CC	for agonists and antagonists by binding and observing the change		
CC	activity. Effective pharmacological agents useful in diagnosis c		
CC	treatment of disease are also identified using APO4 polypeptides		
CC	fragments and APO4 signal transducer molecules that specifically		
CC	with a cytoplasmic domain of APO4 and detecting a change in leve		

06:25:10 2004

us-09-245-198a-1.rng

42.7%; Score 498.8; DB 4; Length 898;
arity 87.0%; Pred No 1.4e-124;
onservative 0; Mismatches 82; Indels 0; Gaps 0;

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|||||
ACCGCCGGAGCCCCCTGAAGTCTCCACAGAGAGAAAGCCAGGATGTGTA 175
|||||
AGGACAGGACCCGTCGGAACCTGAATCCCCACAGAGAAAGCCAGGATCCTGG 369
|||||
TCTTGGAAACAAGTCTCGGCTCGAAGAGTCTCTTAAAGCCGGHAGGCGGG 235
|||||
TCTTGAACCGACTAGTTCGGCTCGCAGAGTGACCTTAAAGCCGGHAAACAGG 429
|||||
GCGGAGCTATTCCAGCCCATTTAGAGTTTCATCTCGGCCAGGACAGGATGGAGCA 295
|||||
GAAAGAGCATCGAGCCCATTTATGAAGTTTCATCCAGACCTGGACAGGACGAGCG 489
|||||
CAGGTGTGGATGGGACAGTGTGGCTGGAGAGACCAAAATCAACAGCTCCAGC 355
|||||
CAGGTGTGGACGGGACAGTGTGGCTGGAGAGACCAAAATCAACAGCTCCAGC 549
|||||
TGGGCTACAGCCGAGATTTGAGGAAATTTACAGTCTCAGGCTGGGCTCTACTAC 415
|||||
TGGGCTACAGCCGAGATTTGAGGAAATTTATAGTCAACCGGCTGGGCTCTACTAC 609
|||||
TCTGTGAGTGTGACTTTGATGAGGAAAGGCTGTCTACTGAGCTGGACTTGTCTG 475
|||||
TCTGTGAGTGTGACTTTGATGAGGAAAGGCTGTCTACTGAGCTGGACTTGTCTG 669
|||||
ACGGTGTGCTGGCCCTGGCTGCTGGAGAAATTTCTCAGCCACAGCAGCAAGCTCT 535
|||||
ATGGTGTGCTGGCCCTGGCTGCTGGAGAAATTTCTCAGCCACTGGGCCAGTTCC 729
|||||
AGCCCCAGCTCCGTTTGTGCCAGGTGTCTGGGCTGTTCGGCTGGGCCAGGGTCT 595
|||||
AGCCCCAGCTCCGCTCTGCCAGGTGTCTGGGCTGTTCGGCTGGGCCAGGGTCC 789
|||||
TTCGGATCCGACCCCTCCGCTGGGCTCATCTTAAGGCTGCCCCCTTCTTAACCTAC 655
|||||
TTCGGATCCGACCCCTCCGCTGGGCTCATCTTAAGGCTGCCCCCTTCTTAACCTAC 849
|||||
GACTCTTTCAAGTTCACTGAGGGCC 685
|||||
GACTCTTCCAGTTCACTGAGGGCC 879

April 7, 2004, 21:32:17
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n search, using sw model

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09-245-198A-2
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SUM62

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681 seqs, 52070155 residues

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ximum Match 100%

-sting first 45 summaries

visProt_42: *

the number of results predicted by chance to have a
s than or equal to the score of the result being printed,
ed by analysis of the total score distribution.

SUMMARIES

seq	ch	Length	DB	ID	Description
0.0	225	1	TN12	MOUSE	O54907 mus musculus
7.8	249	1	TN12	HUMAN	O43508 homo sapien
9.3	272	1	TNF5	CHICK	Q918d8 gallus gall
3.8	316	1	TN11	MOUSE	O35235 m tumor nec
7.8	260	1	TNF5	CANFA	O97626 canis famil
7.8	318	1	TN11	RAT	O9ese2 r tumor nec
7.6	532	1	PBN	HUMAN	P10696 homo sapien
7.6	3664	1	MINT	HUMAN	Q96t58 homo sapien
7.5	244	1	TNFC	HUMAN	Q06643 homo sapien
7.5	244	1	TNFC	PANTR	Q86227 pan troglod
7.5	261	1	TNF5	AOTTR	Q9bdm3 aotus trivi
7.5	261	1	TNF5	CALJA	Q9bnd3 callithrix
7.4	240	1	TN14	HUMAN	O43557 homo sapien
7.4	1237	1	B3A2	MOUSE	P13808 mus musculus
7.4	261	1	TNF5	HUMAN	P29965 homo sapien
7.4	261	1	TNF5	MACMU	Q9bdc7 macaca mula
7.3	240	1	TNF5	MACNE	Q9bdm7 macaca neme
7.3	1237	1	B3A2	RABIT	P48746 oryctolagus
7.2	278	1	TNF6	RAT	P36940 rattus norv
7.2	241	1	TN13	MOUSE	Q9d777 mus musculus
7.2	250	1	TN13	MACMU	Q9xt47 macropus eu
7.1	400	1	TRPB	CHROMO	Q7nud8 chromobacte
7.1	2779	1	LVA	BROME	Q8msa1 drosophila
7.1	535	1	PBI	HUMAN	P05187 homo sapien
6.9	246	1	CIQC	MOUSE	Q02105 mus musculus
6.9	1234	1	B3A2	RAT	P23347 rattus norv
6.9	1465	1	DPQ3	STRMU	Q8dwe0 streptococc
6.8	197	1	TNFB	RABIT	P10154 oryctolagus
6.8	300	1	NTH1	MOUSE	Q35980 mus musculus
6.8	920	1	PARC	SYN3	P73077 synecocyst
6.8	1584	1	U104	CABEL	P23678 caenorhabdi
6.8	2468	1	MAPB	HUMAN	P46821 homo sapien
6.8	260	1	TNF5	FELCA	O97605 felis silve

34	78.5	6.8	480	1	KCG2_RAT	Q9qr03 rattu
35	78.5	6.8	1164	1	PHYD_ARATH	P42497 arab:
36	78	6.7	285	1	T13B_HUMAN	Q9Y275 homo
37	78	6.7	495	1	GATB_METAC	Q8thj0 methi
38	78	6.7	495	1	GATB_METAC	Q8px10 methi
39	78	6.7	763	1	APP2_HUMAN	Q06481 homo
40	77.5	6.7	201	1	TNFB_MACMU	Q9xt48 macr
41	77.5	6.7	261	1	TNFB_PIG	Q95mq5 sus
42	77	6.6	279	1	TNF6_MOUSE	P41047 mus
43	77	6.6	817	1	NAH1_BOVIN	Q28036 bos
44	76.5	6.6	214	1	SMP_ECOLI	P18838 eschu
45	76.5	6.6	788	1	NASP_HUMAN	P49321 homo

ALIGNMENTS

RESULT 1	TN12_MOUSE	STANDARD;	PRT;	225 AA.
ID	AC	O54907; Q9CTP2;		
DT	DT	28-FEB-2003 (Rel. 41, Created)		
DT	DT	28-FEB-2003 (Rel. 41, Last sequence update)		
DT	DT	28-FEB-2003 (Rel. 41, Last annotation update)		
DE	DE	tumor necrosis factor ligand superfamily member 12 (TNF-related w		
DE	DE	inducer of apoptosis) (TWEAK) (fragment).		
GN	GN	TNFSF12.		
OS	OS	Mus musculus (Mouse).		
OC	OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	OC	Mammalia; Eutharia; Rodentia; Sciurognathi; Muridae; Murinae; Mus		
OX	OX	NCBI_TaxID=10090;		
LN	LN	[1]		
RP	RP	SEQUENCE FROM N.A.		
RC	RC	TISSUE=Peritoneal macrophage;		
RX	RX	MEDLINE=98070415; PubMed=9405449;		
RA	RA	Chicheportiche Y., Bourdon P.R., Xu H., Hsu Y.-M., Scott H.,		
RA	RA	Hession C., Garcia I., Browning J.L.;		
RT	RT	"TWEAK, a new secreted ligand in the tumor necrosis factor family		
RT	RT	weakly induces apoptosis."		
RL	RL	J. Biol. Chem. 272:32401-32410(1997).		
LN	LN	[2]		
RP	RP	SEQUENCE OF 83-225 FROM N.A.		
RC	RC	STRAIN=C57BL/6J; TISSUE=Retina;		
RX	RX	MEDLINE=21085660; PubMed=11217851;		
RA	RA	Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.		
RA	RA	Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.		
RA	RA	Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamataka I.		
RA	RA	Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.		
RA	RA	Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,		
RA	RA	Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,		
RA	RA	Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush		
RA	RA	Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Barsh G.,		
RA	RA	Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., de Bonaldo M.F.,		
RA	RA	Blake J., Boffelli D., Bojunga N., Carninci P., Fujita M., Gariboldi M.,		
RA	RA	Brownstein M.J., Bult C., Fletcher C., Hume D.A., Kamiya M., Lee N.H		
RA	RA	Gustincich S., Hill D., Hofmann M., Mazzarelli J., Mombarts P.,		
RA	RA	Lyons P., Marchionni L., Mashima J., Rodriguez I., Sakamoto N.,		
RA	RA	Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,		
RA	RA	Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.		
RA	RA	Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilmir		
RA	RA	Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.		
RA	RA	Hayashizaki Y.;		
RT	RT	"Functional annotation of a full-length mouse cDNA collection.";		
RL	RL	Nature 409:685-690(2001).		
CC	CC	-!- FUNCTION: Binds to FN14 and possibly also to TNFRSF12/APO3. A		
CC	CC	inducer of apoptosis in some cell types. Promotes angiogenesis		
CC	CC	the proliferation of endothelial cells. Mediates NF-KappaB		
CC	CC	activation (by similarity).		
CC	CC	-!- SUBUNIT: Homotrimer (Potential).		
CC	CC	-!- SUBCELLULAR LOCATION: Type II membrane protein and secreted		
CC	CC	similarity).		
CC	CC	-!- TISSUE SPECIFICITY: Widely expressed.		
CC	CC	-!- PTM: The soluble form is produced from the membrane form by		

ic processing (By similarity).
Y: Belongs to the tumor necrosis factor family.

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10: AAC53517.1; -
19: BAB32249.1; -
1259: Tnfef12.
1006052: TNF family.
1008983: TNF-like.
1: TNF; 1.
17: TNF; 1.
1251: TNF 1; FALSE_NEG.
1049: TNF 2; 1.
Hogensis; Apoptosis; Transmembrane; Glycoprotein;
1: 1
1 225 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 12, MEMBRANE FORM.
70 225 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 12, SECRETED FORM (BY SIMILARITY).
1 21 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
(POTENTIAL).
22 225 EXTRACELLULAR (POTENTIAL).
69 70 CLEAVAGE (BY SIMILARITY).
15 115 N-LINKED (GLCNAC... (POTENTIAL).
15 AA: 24781 MW; 90CA12CC0480659B CRC64;
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Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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VPRPSAPKGRARRAIAAHYVHPRPGDGAQAGVDGTVSGWEEKINSSSL 120
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IQIGFTVIRAGLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLEFSAATASSPG 180
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RLCQVSGLLPLRPGSSLRIRTLPAHLKAAPFLTYGLFQVH 225

STANDARD; PRT; 249 AA.
27;
(Rel. 41, Created)
(Rel. 41, Last sequence update)
(Rel. 42, Last annotation update)
is factor ligand superfamily member 12 (TNF-related weak
potosis) (TWEAK) (APO3 ligand).
PO3L OR DR3LG.
(Human).
etazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Primates; Catarrhini; Hominidae; Homo.
506;
M N.A., AND N-TERMINUS OF SOLUBLE FORM.
liver, and Tonsil;

RX MEDLINE=98070415; PubMed=9405449;
RA Chicheportiche Y., Bourdon P.R., Xu H., Hsu Y.-M., Scott H.,
RA Hession C., Garcia I., Browning J.L.;
RT "TWEAK, a new secreted ligand in the tumor necrosis factor family
weakly induces apoptosis.";
RL J. Biol. Chem. 272:32401-32410(1997).
[2]
RP SEQUENCE FROM N.A.
RC TISSUE=Fetal kidney;
RX MEDLINE=98228355; PubMed=9560343;
RA Marsters S.A., Sheridan J.F., Pitti R.M., Brush J., Goddard A.,
RA Ashkenazi A.;
RT "Identification of a ligand for the death-domain-containing recep
Ap03.";
RL Curr. Biol. 8:525-528(1998).
[3]
RP SEQUENCE FROM N.A.
RC TISSUE=Tonsil;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins P.S., Wagner L., Shenmen C.M., Schuler G.I.
RA Altshul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz
RA Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.I
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A.C., Rodrigues S., Sanchez
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickinson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
[4]
RN FUNCTION.
RP MEDLINE=99185061; PubMed=10085077;
RA Lynch C.N., Wang Y.C., Lund J.K., Chen Y.-W., Leal J.A., Wiley S
RT "TWEAK induces angiogenesis and proliferation of endothelial cel
J. Biol. Chem. 274:8455-8459(1999).
CC -!- FUNCTION: Binds to FN14 and possibly also to TNFRSF12/AP03. I
inducer of apoptosis in some cell types. Mediates NF-kappaB
activation. May promote angiogenesis and the proliferation o
endothelial cells.
CC -!- SUBUNIT: Homotrimer (Potential).
CC -!- SUBCELLULAR LOCATION: Type II membrane protein and secreted.
CC -!- TISSUE SPECIFICITY: Highly expressed in adult heart, pancrea
skeletal muscle, brain, colon, small intestine, lung, ovary,
prostate, spleen, lymph node, appendix and peripheral blood
lymphocytes. Low expression in kidney, testis, liver, placen
thymus and bone marrow. Also detected in fetal kidney, liver
lung and brain.
CC -!- PTM: The soluble form derives from the membrane form
by proteolytic processing.
CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
CC -!- CAUTION: Ref.3 sequence differs from that shown due to a
frameshift in position 125.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a col
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the European Bioinformatics Institute. There are no restrictio
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modified and this statement is not removed. Usage by and for
entities requires a license agreement (See <http://www.isb-sib.ch>
or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF030099; AAC51923.1; -
CC EMBL; AF055872; AAC39724.1; -
DR
DR

```

; AAH19047.1; ALT_FRAME.
; 927; TNFSF12.
; C: integral to plasma membrane; TAS.
; F: receptor binding; TAS.
; P: induction of apoptosis; TAS.
; S: signal transduction; TAS.
06052; TNF family.
08983; TNF-like.
; TNF; 1.
; TNF; 1.
; 51; TNF 1; FALSE_NEG.
149; TNF 2; 1.
ogenesis; Apoptosis; Transmembrane; Glycoprotein;

1 249 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 12, MEMBRANE FORM.
14 249 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 12, SECRETED FORM.
1 21 CYTOPLASMIC (POTENTIAL).
22 42 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
(POTENTIAL).
43 249 EXTRACELLULAR (POTENTIAL).
33 94 CLEAVAGE.
39 139 N-LINKED (GLCNAC. . .).
3 AA; 27216 MW; E660843361C28EBA CRC64;

87.8%; Score 1020; DB 1; Length 249;
arity 88.8%; Pred. No. 9.4e-83;
conservative 9; Mismatches 16; Indels 0; Gaps 0;

LALACGLLVVSLGSWATLSAQPSOELTAEDRRPELNPQTEESQDVVPFL 61
LALACGLLVVSLGSRASLSQAEPAQELVAEDQDPSLNPQTEESQDPAPFL 85
RPRRSAPKGRKARRAIAAHYEVHPRPGDGAQAGVDGTVSGWEERINSSSPLR 145
IGFTVIRAGLYLYCQVHFDEGKAVYKLDLLVGVLAIRCLLEFSATASPGP 181
IGEFIVTRAGLYLYCQVHFDEGKAVYKLDLLVGVLAIRCLLEFSATASLIGP 205
QCVSGLLAPRGSSLRITLPWAHLKAAPFLTYFGLFQVH 225
QCVSGLLAPRGSSLRITLPWAHLKAAPFLTYFGLFQVH 249

STANDARD; PRT; 272 AA.

Rel. 41, Created
Rel. 43, Last sequence update
Rel. 43, Last annotation update
s factor ligand superfamily member 5 (CD40 ligand) (CD40-
tein)
OLG OR CD40L.
(Chicken).
tazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;

31;
(N.A.
leghorn; TISSUE=Spleen;
Young J.R., Burnside J.;
putative chicken CD40 ligand";
P-2003) to the EMBL/GenBank/DBJ databases.
Cytokine that binds to TNFRSF5. Mediates B-cell
ation in the absence of co-stimulus as well as IgE
n in the presence of IL-4. Involved in immunoglobulin
tching (By similarity).

```

```

CC -!- SUBUNIT: Homotrimer (By similarity).
CC -!- SUBCELLULAR LOCATION: Type II membrane protein. Also exists a
CC extracellular soluble form (By similarity).
CC -!- PTM: The soluble form derives from the membrane form by
CC proteolytic processing (By similarity).
CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
CC
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CC entities requires a license agreement (See http://www.isb-sib.ch/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AJ243435; CAB95748.2; -.
CC HSP; P29965; ITALY.
CC GO; GO:0016021; C: integral to membrane; ISS.
CC GO; GO:0005174; F: CD40 receptor binding; ISS.
CC GO; GO:0042100; P: B-cell proliferation; ISS.
CC GO; GO:0006954; P: inflammatory response; ISS.
CC GO; GO:0007159; P: leukocyte cell adhesion; ISS.
CC GO; GO:0030168; P: platelet activation; ISS.
CC InterPro; IPR003263; TNF_5.
CC InterPro; IPR006052; TNF family.
CC InterPro; IPR008983; TNF-like.
CC InterPro; IPR003636; TNF_subf.
CC Pfam; PF00229; TNF; 1.
CC PRINTS; PR01702; CD40LIGAND.
CC ProDom; PD008600; TNF_5; 1.
CC ProDom; PD002012; TNF_subf; 1.
CC SMART; SM00207; TNF; 1.
CC PROSITE; PS00251; TNF 1; 1.
CC PROSITE; PS00049; TNF 2; 1.
CC Cytokine; Transmembrane; Glycoprotein; Signal-anchor.
CC CHAIN 1 272
CC CHAIN 111 272
CC DOMAIN 1 23
CC TRANSMEM 24 44
CC DOMAIN 45 272
CC SITE 110 111
CC DISULFID 190 229
CC CARBOHYD 124 124
CC CARBOHYD 146 146
CC CARBOHYD 251 251
CC SEQUENCE 272 AA; 30832 MW; 8CD0338A924E044B CRC64;

Query Match 9.3%; Score 108.5; DB 1; Length 272;
Best Local Similarity 22.5%; Pred. No. 0.024;
Matches 58; Conservative 40; Mismatches 107; Indels 53; G

Qy 1 VLSGLGLALGILLVSVLSGSWATLSAQ-----FPSQELTAEDRRPE-----
Db 34 VQTIGTVLFCLYLHKMKDKMEVLSDNEDYIFLRKVKQCTGDEQKSTLLDCEKVI
Qy 46 ELNPQTEESQDVVPFLQLVPRPSAPGRK-----ARPRRAIAAHYEVHPRPGOI
Db 94 DLQCKORTASEELPKFEMHGRGHEPHLKSRNETSVAEKRPPIATHLA-----
Qy 101 GVDGTGVSGWEETK-INSSSPLRYDRQIGCEFTVIRAGLYLYCQVHFDEGKA-----
Db 146 NTVTRVLKMTTSTYAPTSSLSIYHE--GKLKVEKAGLYIYSQVSFCTKAAASAPI
Qy 152 YLKLDELVLVGVLAIRCLLEFSATASPGPQLRCQV-----SGLLPLRPGSSLR.
Db 204 YLYLPMEDRL-LMKGLDTHSTSTA-----LCELOSIREGGVFLRQGDVMVF
Qy 207 WAHLKAAPFLTYFGLFQV 224
Db 255 STAVNVNPGNTYFGMKL 272

```

STANDARD; PRT; 316 AA.
 06; Q9JJK8; Q9JJK9; Q9R1Y0;
 (Rel. 40, Created)
 (Rel. 40, Last sequence update)
 (Rel. 42, Last annotation update)
 is factor ligand superfamily member 11 (Receptor activator
 actor kappa B ligand) (RANKL) (TNF-related activation-
 ion factor) (TRANCE) (Osteoprotegerin ligand) (OPGL) (Osteoclast
 ion factor) (ODF) (Osteoclastogenesis-inhibitory factor)
 ANKL OR TRANCE OR OPGL.
 (Mouse).
 stazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Theria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 3090;
 M N.A. (ISOFORM 1).
 3112; PubMed=9312132;
 3112; Arron J., Robinson E., Orlicki J., Chao M.,
 ., Cayani E., Bartlett F.S. III, Frankel W.N., Lee S.Y.,
 novel ligand of the tumor necrosis factor receptor family
 as C-Jun N-terminal kinase in T cells.";
 n. 272:25190-25194(1997).
 M N.A. (ISOFORM 1).
 c lymphoma;
 2977; PubMed=9367155;
 ., Maraskovsky E., Billingsley W.L., Dougall W.C.,
 ., Roux E.R., Teepe M.C., Dubose R.F., Cosman D.,
 of the TNF receptor and its ligand enhance T-cell growth
 c-cell function.";
 75-179(1997).
 M N.A. (ISOFORM 1).
 marrow;
 7661; PubMed=9568710;
 Timms E., Tan H.-L., Kelley M.J., Dunstan C.R.,
 Elliott R., Colombero A., Elliott G., Scully S., Hsu H.,
 Hawkins N., Davy E., Capparelli C., Eli A., Qian Y.-X.,
 Sarosi I., Shalhoub V., Senaldi G., Guo J., Delaney J.,
 erin ligand is a cytokine that regulates osteoclast
 ion and activation.";
 176(1998).
 M N.A. (ISOFORM 1).
 marrow stroma;
 8248; PubMed=9520411;
 hima N., Nakagawa N., Yamaguchi K., Kinoshita M.,
 -I., Tomoyasu A., Yano K., Goto M., Murakami A., Tsuda E.,
 Higashio K., Udagawa N., Takahashi N., Suda T.;
 differentiation factor is a ligand for
 rin/osteoclastogenesis-inhibitory factor and is identical
 NK1.";
 Acad. Sci. U.S.A. 95:3597-3602(1998).
 M N.A. (ISOFORM 1).
 4075; PubMed=10196481;
 Kodaira K., Mizuno A., Yasuda H., Shima N., Murakami A.,
 (characterization of the gene encoding mouse osteoclast
 ion factor.";
 -127(1999).
 M N.A. (ISOFORMS 1; 2 AND 3).
 RX MEDLINE=21150053; PubMed=11250921;
 Ikeda T., Kasai M., Utsuyama M., Hirokawa K.;
 "Determination of three isoforms of the receptor activator of nu
 factor-kappaB ligand and their differential expression in bone a
 thymus.";
 RL Endocrinology 142:1419-1426(2001).
 [7]
 RN SEQUENCE OF 139-147, PROCESSING, AND N-GLYCOSYLATION.
 RP MEDLINE=99240759; PubMed=10224132;
 RX Lum L., Wong B.R., Josien R., Becherer J.D., Erdjument-Bromage H
 RA Schindler J., Tempst P., Choi Y., Blobel C.P.;
 "Evidence for a role of a tumor necrosis factor-alpha
 (TNF-alpha)-converting enzyme-like protease in shedding of TRANC
 TNF family member involved in osteoclastogenesis and dendritic c
 survival.";
 RT J. Biol. Chem. 274:13613-13618(1999).
 RL [8]
 RN X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS) OF 158-316.
 RP MEDLINE=21464816; PubMed=11581298;
 RX Lam J., Nelson C.A., Ross F.P., Teitelbaum S.L., Fremont D.H.;
 RA "Crystal structure of the TRANCE/RANKL cytokine reveals determin
 RT of receptor-ligand specificity.";
 RL J. Clin. Invest. 108:971-979(2001).
 [9]
 RN X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 137-316.
 RP MEDLINE=21839021; PubMed=11733492;
 RX Ito S., Wakabayashi K., Ubukata O., Hayashi S., Okada F., Hata T
 RA "Crystal structure of the extracellular domain of mouse RANK lig
 RT 2.2-A resolution.";
 RL J. Biol. Chem. 277:6631-6636(2002).
 CC -!- FUNCTION: Cytokine that binds to TNFRSF1B/OPG and to
 CC TNFRSF1A/RANK. Osteoclast differentiation and activation fa
 CC Augments the ability of dendritic cells to stimulate naive T
 CC proliferation. May be an important regulator of interactions
 CC between T cells and dendritic cells and may play a role in t
 CC regulation of the T cell-dependent immune response. May also
 CC an important role in enhanced bone-resorption in humoral
 CC hypercalcemia of malignancy.
 CC -!- SUBUNIT: Homotrimer.
 CC -!- SUBCELLULAR LOCATION: Type II membrane protein and secreted
 CC (isoforms 1 and 2); Cytoplasmic (isoform 3).
 CC -!- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=3;
 CC Name=1;
 CC IsoId=O35235-1; Sequence=Displayed;
 CC Name=2;
 CC IsoId=O35235-2; Sequence=VSP_006449;
 CC Name=3;
 CC IsoId=O35235-3; Sequence=VSP_006448;
 CC -!- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN THYMUS AND LYMPH NOD
 CC NOT IN NONLYMPHOID TISSUES AND IS ABUNDANTLY EXPRESSED IN T
 CC BUT NOT IN B CELLS. A HIGH LEVEL EXPRESSION IS ALSO SEEN IN
 CC TRABECULAR BONE AND LUNG.
 CC -!- PTM: N-glycosylated.
 CC -!- PTM: The soluble form of isoform 1 derives from the membrane
 CC by proteolytic processing. The cleavage may be catalyzed by
 CC ADAM17. A further shorter soluble form was observed.
 CC -!- DISBASE: Deficiency in TNFSF11 results in failure to form lc
 CC alveolar mammary structures during pregnancy, resulting in c
 CC of newborns. Trance-deficient mice show severe osteopetrosis
 CC no osteoclasts, marrow spaces, or tooth eruption, and exhibi
 CC profound growth retardation at several skeletal sites, inclu
 CC the limbs, skull, and vertebrae and have marked chondrodyspl
 CC with thick, irregular growth plates and a relative increase
 CC hypertrophic chondrocytes.
 CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
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AAC71061.1; -
AAC86812.1; -
AAC40113.1; -
BAA25425.1; -
BAA36970.1; -
BAA36970.1; JOINED.
BAA36970.1; JOINED.
BAA36970.1; JOINED.
BAA97257.1; -
BAA97258.1; -
BAA97259.1; -
BAA97259.1; -
JAN-03.
JAN-03.
J89; Tnfsf11.
P: Organogenesis; IMP.
P: Ossification; IMP.
006052; TNF family.
008983; TNF_like.
003636; TNF_subf.
; TNF; 1.
012; TNF_subf; 1.
7; TNF; 1.
251; TNF; 1; FALSE_NEG.
049; TNF; 2; 1.
fermentation; Receptor; Glycoprotein; Transmembrane;
3D-structure; Alternative splicing.
1 316 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 11, MEMBRANE FORM.
39 316 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 11, SOLUBLE FORM.
1 48 CYTOPLASMIC (POTENTIAL).
49 69 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
(POTENTIAL).
EXTRACELLULAR (POTENTIAL).
70 316 N-LINKED (GLCNAC. . .) (POTENTIAL).
38 139 N-LINKED (GLCNAC. . .) (POTENTIAL).
97 197 N-LINKED (GLCNAC. . .) (POTENTIAL).
262 62 Missing (in isoform 3).
1 117 /FTid=VSP_006448.
14 44 /FTid=VSP_006449.
isoform 2).
99 99 G -> D (IN REF. 2).
41 143 MISSING (IN REF. 5).
143 169
64 169
71 172
81 182
86 187
91 192
94 196
98 201
203 203
204 207
224 224
225 227
245 245
8.8%; Score 102.5; DB 1; Length 316;
larity 24.3%; Pred. No. 0.096;
Conservative 37; Mismatches 107; Indels 65; Gaps 13;
ALACILGLLVVYVSLGSMATLSAQ-EPSELTAEDR-----REPPELNPQT 51
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
ALLGLGLGVVCSIALFLYFRAQMDPNR---ISEDSTHCFYRLRLHENAGLQDST 104
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QDVVP-----FLEQLVRRR--SAPK-----GRKAPRRRAIA 84
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
EDTLPSDCRMKQAFQAGVQKELQHVGVGRFSGAPAMMEGSLWLDVAQRGFEAPF 164
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
EVHPRQDGAQGVDTGVSGEE-----TKINSSSFLRYDRQIGFTVIRAGLYLY 140
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

```


SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)

(POTENTIAL).

EXTRACELLULAR (POTENTIAL) .

CLEAVAGE (BY SIMILARITY).

POTENTIAL.

N-LINKED (GLCNAC...) (POTENTIAL) .

60 AA; 28688 MW; 604F69A19E9EB70 CRC64;

7.8%; Score 90.5; DB 1; Length 260;

arity 28.1%; Pred.No.0.96;

conservative 16; Mismatches 53; Indels 31; Gaps 5;

NSSSPLRYDRQGTGEFT-----VIRAGLYLVCQHVFDEGKAVYLKL 155
| | : | : ||||| :
| : | : | | : | : | : | :
NPASVLWRWAP-KGYTITISSNLVSLENGKQLAVKRQGIIYVAQTFCSNRAASSQA 186

TGVLALRCLIEFSAT-----AASSPPQLRLCQVS---GILLPRPGSSLRIRTL 205
| | | | | ||||| :
----VASLCIHSFGTSRVLLRAASSRGSSXPCQQSIHLGGVFEIHLPKGASFVNVT 243

ELKAAPFLTYFGLFPQV 224
| : | : | :
| VSHGTGTFSGLLKL 260

STANDARD; PRT; 318 AA.

[Rel. 41, Created]
[Rel. 41, Last sequence update)
[Rel. 41, last annotation update)
s factor ligand superfamily member 11 (Receptor activator
factor kappa B ligand) (RANKL) (TNF-related activation-
line) (TRANCE) (Osteoprotegerin ligand) (OPGL) (Osteoclast
on factor) (ODF).
ANKL OR TRANCE OR OPGL.
jicus (Rat).
starcia; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
1116;

4 N.A.
l bone;
J.K., Huang L., Gao X.H., Laird R., Liu D., Wysocki S.,
ence and functional characterization of the rat
receptor activator of NF-kB ligand." ;
c. Res. 15:2178-2186(2000).

266-318 FROM N.A.
ar 344;
2371; PubMed=11804028;
Kim N., van Wesenbeeck L., Mackay C., Mason-Savas A.,
Popoff S.N., Lengner C., van-Hul W., Choi Y.,
r.;
at the rat osteoporotic mutation toothless (tl) is not in
(TRANCE), RANKL, ODF, OPGL) gene.";
Biol. 45:853-859(2001).
: Cytokine that binds to TNFRSF11B/OPG and to
A/RANK. Osteoclast differentiation and activation factor.
the ability of dendritic cells to stimulate naive T-cell
ation. May be an important regulator of interactions
T cells and dendritic cells and may play a role in the
on of the T cell-dependent immune response. May also play
tant role in enhanced bone-resorption in humoral
cemia of malignancy.
Homotrimer (By similarity).
LAR LOCATION: Type II membrane protein and secreted (By
ty).
PECIFICITY: Highly expressed in thymus and bone tissues.
soluble form derives from the membrane form by

Human).
azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Sirta; Primates; Catarrhini; Hominidae; Homo.

N.A.
79; PubMed=2162249;
aus A.W.;
a Nagao-type, phosphatidylinositol-glycan anchored
hataase in human choriocarcinomas";
:3956-3962(1990).

N.A.
96; PubMed=2745460;
atanabe T., Li W.L., Soong B.-W., Chou J.Y.;
the germ cell alkaline phosphatase gene in human
a cells";
264:12611-12619(1989).

N.A.
32; PubMed=2834730;
anes T.;
ved Nagao isozyme is encoded by a germ-cell alkaline
ne.";
ad. Sci. U.S.A. 85:3024-3028(1988).

N.A.
11; PubMed=2297757;
s J.W., Sack T.L., Kim Y.S.;
ning of complementary DNAs encoding alkaline
human colon cancer cells.";
:1085-1091(1990).

N.A.
a;
57; PubMed=12477932;
Feingold E.A., Grouse L.H., Derge J.G.,
Collins F.S., Wagner L., Shemmen C.W., Schuler G.D.,
Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
Marusina K., Farmer A.A., Rubin G.M., Hong L.,
Soares M.B., Bonaldi M.F., Casavant T.L., Scheetz T.E.,
Usdin T.B., Tohiyuki S., Carninci P., Prange C.C.,
Mellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
Ewan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
Orley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
on E., Ketterman M., Madan A., Rodriguez S., Sanchez A.,
dan A., Young A.C., Shevchenko Y., Bouffard G.G.,
Touchman J.W., Green E.D., Dickson M.C.,
Grimwood J., Schmutz J., Myers R.M.,
S.N., Krzywinski M.I., Skaleka U., Smallos D.E.,
chein J.E., Jones S.J.M., Marra M.A.;
id initial analysis of more than 15,000 full-length
se cDNA sequences";
ad. Sci. U.S.A. 99:16999-16903(2002).

157 FROM N.A.
578; PubMed=3387245;
a H., Kan Y.W., Kam W.;
a sequence of a putative human placental alkaline
like gene";

Res. 16:5694-5694(1988).
ACTIVITY: An orthophosphoric monoester + H(2)O = an
phosphate.
monomer.

AR LOCATION: Attached to the membrane by a GPI-anchor.
SPECIFICITY: TRACE AMOUNTS IN THE TESTIS AND THYMUS,
EVATED AMOUNTS IN GERM CELL TUMORS.
BOUS: In most mammals there are four different isozymes:
, placental-like, intestinal and tissue non-specific

CC (liver/bone/kidney).
CC -!- SIMILARITY: Belongs to the alkaline phosphatase family.
CC
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CC EMBL; X55958; CAA39425.1; -
CC EMBL; J04948; AAA51700.1; -
CC EMBL; J03252; AAA98616.1; -
CC EMBL; X53279; CAA37374.1; -
CC EMBL; BC014139; AAH14139.1; -
CC EMBL; X07247; CAA30232.1; ALT_SEQ.
CC PIR; S12076; S12076.
CC DR HSSP; P00634; LAJC.
CC DR Siena-2DPAGE; P10696; -
CC DR Genew; HGNC:441; ALPPL2.
CC MIM; 171810; -

CC GO; GO:0016020; C:membrane; NAS.
CC GO; GO:0004035; F:alkaline phosphatase activity; NAS.
CC GO; GO:0016310; P:phosphorylation; NAS.
CC InterPro; IPR001952; Alk phosphatse.
CC Pfam; PF0245; alk phosphatase; 1.
CC PRINTS; PR00113; ALKPHPTASE.
CC ProDom; PD001868; Alk_phosphatse; 1.
CC SMART; SM00090; alkPFC; 1.
CC PROSITE; PS00123; ALKALINE_PHOSPHATASE; 1.
CC HydroLase; Zinc; Magnesium; Phosphorylation; Transmembrane;
CC Multigene family; Glycoprotein; GPI-anchor; Signal.
CC SIGNAL; 19
CC POTENTIAL.

CC CHAIN; 20 503
CC PROPEP; 504 532
CC DISULFID; 140 202
CC DISULFID; 486 493
CC ACT_SITE; 111 111
CC CARBOHYD; 141 141
CC CARBOHYD; 268 268
CC CONFLICT; 57 57
CC CONFLICT; 152 152
CC CONFLICT; 178 178
CC CONFLICT; 260 260
CC CONFLICT; 273 273
CC CONFLICT; 316 316
CC CONFLICT; 380 380
CC CONFLICT; 498 498
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CC CONFLICT; 531 531
CC SEQUENCE; 532 AA; 57315 MW; 84AB5B28F13D6D82 CRC64;

CC ALKALINE PHOSPHATASE, PLACENTAL-LIKE
CC REMOVED IN MATURE FORM.
CC BY SIMILARITY.
CC BY SIMILARITY.
CC PHOSPHOSERINE INTERMEDIATE.
CC N-LINKED (GLCNAC. . .) (POTENTIAL).
CC N-LINKED (GLCNAC. . .) (POTENTIAL).
CC I -> M (IN REF. 3).
CC V -> M (IN REF. 2, 4, 5 AND 6).
CC T -> A (IN REF. 4 AND 5).
CC H -> R (IN REF. 5).
CC L -> M (IN REF. 4 AND 5).
CC L -> R (IN REF. 1, 4 AND 5).
CC V -> L (IN REF. 2).
CC P -> S (IN REF. 4).
CC A -> T (IN REF. 4).

CC Query Match 7.6%; Score 88; DB 1; Length 532;
CC Best Local Similarity 28.6%; Pred. No. 3.4;
CC Matches 44; Conservative 16; Mismatches 44; Indels 50; G;
QY 76 KAPRRAIAAHYEV---HPRPG---ODGQAQAGVDGTGSGWEETKINSSSPRYDRQ
Db 400 KARDKA---YTVLLYNGPGVVLKDGARPDVTSESGSPYRQQAAYPLDGETH
QY 130 TVIRAGLYLYYQVHFDEKAVYLKDLLVNGV-----LALRCLLEFSA---
Db 456 VAV-----FARGPOAH-----LVHGVQEQTFIAHVMAFAACLEPYTACD;
QY 174 ---TAASSPGQLRLCQVSGLLPLRPGSSIRINT 204
Db 499 AGTTDAHPGPSV-----VPALPLLAGTLLLT 528

CC RESULT 8

CC MINT_HUMAN

CC ID MINT_HUMAN STANDARD; PRT; 3664 AA.


```

34; TNECROSISFCT.
:012; TNF subf; 1.
17; TNF; 1.
1251; TNF_1; 1.
1049; TNF_2; 1.
membrane; Glycoprotein; Signal-anchor;
splicing; Polymorphism.
1 18 CYTOPLASMIC (POTENTIAL).
19 48 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
(POTENTIAL).
49 244 EXTRACELLULAR (POTENTIAL).
222 222 N-LINKED (GLCNAC...;) (POTENTIAL).
53 77 GLVTETADPGAQOGLGFKLPEE -> GLGFRSCQRRSQ
KQISAPGSQLPTS (in isoform 2).
/FTId-VSP 056441.
Missing (in isoform 2).
78 244 /FTId-VSP_006442.
70 70 G->E.
/FTId-VAR_013025.
84 84 S->R.
/FTId-VAR_016331.
87 87 L->F.
/FTId-VAR_016332.
11 111 A->P.
/FTId-VAR_013026.
60 69 DPGAQOGL -> GLSAPGSGRT (IN REF. 2;
AAB37342).
14 AA; 25390 MW; F41569459830ED4C CRC64;
7.5%; Score 87; DB 1; Length 244;
arity 23.0%; Pred. No. 1.6;
Conservative 26; Mismatches 80; Indels 108; Gaps 13;
ALA---CLGLLVVSLGSWATLSAQEPSQBELTAEDR-----REPPPEL 47
AVAGATSLVTLIAVPITVLAVLVPDQOGLVTETADPGAQOGLGFKLPEE 77
[PESQDVVPLEQLVPRRSAPKGRKARPRRAIAAHYEVHPREGQDGAQGVDTYS 107
T---DLSPGLP-----AAHLGAPLKGQ-----L 102
ETKINS--SSPLRYDRQIGFETVIRAGLYLYCQVHF-----DEKAVYLKL 155
[TKTEQAFLTSGTQFSDAEG-LALPDQGLYLYCLVGRAPPGGGDPQGRSVTLRS 161
NGVLALRCLEEFSAATAASPG-PQLRL-----CQVS 189
-----YRAGAYCGPTPELLLEGAETVTPVLPDARRQGYGLMYTSVGFG 208
PLRPGSLRIRTLPAHLKAAPFL---TYFGLFQV 224
LRRGERVYVNI---SHPDMVDFAFGKTFPGAVMV 243
STANDARD; PRT; 244 AA.
(Rel. 43, Created)
(Rel. 43, Last sequence update)
(Rel. 43, Last annotation update)
beta (LT-beta) (Tumor necrosis factor C) (TNF-C) (Tumor
tor ligand superfamily member 3).
3 OR TNFC.
tes (Chimpanzee).
atazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Cheria; Primates; Catarrhini; Homnidae; Pan.
598;
N.A.
1002; PubMed=12493009;
Shiina T., Anzai T., Kohara S., Inoko H.;
genomic analysis of the MHC: the evolution of class I

```

RT duplication blocks, diversity and complexity from shark to man."
RN Immunol. Rev. 190:95-122(2002).
RP SEQUENCE FROM N.A.
RX MEDLINE=22709134; PubMed=12799463;
RA Anzai T., Shiina T., Kimura N., Yanagiya K., Kohara S., Shigenar.
RA Yamagata T., Kuleki J.K., Naruse T.K., Fujimori Y., Fukuzumi Y.,
RA Yamazaki M., Tashiro H., Iwamoto C., Umehara Y., Imanishi T.,
RA Meyer A., Ikeo K., Gojobori T., Bahram S., Inoko H.;
RT "Comparative sequencing of human and chimpanzee MHC class I regi.
RT unveils insertions/deletions as the major path to genomic
RT divergence."
RT Proc. Natl. Acad. Sci. U.S.A. 100:7708-7713(2003).
RL -!- FUNCTION: Cytokine that binds to LTB4/TNFRSF3. May play a sp.
CC role in immune response regulation. Provides the membrane an.
CC for the attachment of the heterotrimeric complex to the cell
CC surface (By similarity).
CC -!- SUBUNIT: Heterotrimer of either two LTB and one LTA subunits
CC (less prevalent) two LTA and one LTB subunits (By similarity).
CC -!- SUBCELLULAR LOCATION: Type II membrane protein (Potential).
CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AB054536; BAB3881.1; -.
CC EMBL; AB100082; BAC78156.1; -.
CC InterPro; IPR006053; TNF abc.
CC InterPro; IPR006052; TNF family.
CC InterPro; IPR008983; TNF like.
CC InterPro; IPR003636; TNF_subf.
CC Pfam; PF00229; TNF; 1.
CC PRINTS; PR01234; TNECROSISFCT.
CC ProDom; PD002012; TNF subf; 1.
CC SMART; SM00207; TNF; 1.
CC PROSITE; PS00251; TNF_1; 1.
CC PROSITE; PS50049; TNF_2; 1.
CC Cytokine; Transmembrane; Glycoprotein; Signal-anchor.
KW CYTOPLASMIC (POTENTIAL).
FT DOMAIN 1 18 SIGNAL-ANCHOR (TYPE-II MEMBRANE PRO
FT TRANSMEM 19 48 (POTENTIAL).
FT DOMAIN 49 244 EXTRACELLULAR (POTENTIAL).
FT CARBOHYD 222 222 N-LINKED (GLCNAC...;) (POTENTIAL).
FT SEQUENCE 244 AA; 25420 MW; A4047858335DSB97 CRC64;
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Best Local Similarity 23.0%; Pred. No. 1.6;
Matches 64; Conservative 26; Mismatches 80; Indels 108;
QY 3 SLGLALA---CLGLLVVSLGSWATLSAQEPSQBELTAEDR-----R
DB 18 SLLAVAGATSLVTLIAVPITVLAVLVPDQOGLVTETADPGAQOGLGFGQ
QY 48 NPQTEESQDVVPLEQLVPRRSAPKGRKARPRRAIAAHYEVHPREGQDGAQAGV
DB 78 ERET---DLSPGLP-----AAHLGAPLKGQ-----
QY 108 GWEETKINS--SSPLRYDRQIGFETVIRAGLYLYCQVHF-----DSGKA
DB 103 GWETTKEAFLTSGTQFSDAEG-LALPDQGLYLYCLVGRYGRTPPPGGDPQGRS
QY 156 DLLVNGVLALRCLEEFSAATAASPG-PQLRL-----
DB 162 SL-----YRAGAYCGPTPELLLEGAETVTPVLPDARRQGYGLMYT
QY 190 GLPLRPGSLRIRTLPAHLKAAPFL---TYFGLFQV 224
DB 209 GLVQRRGERVYVNI---SHPDMVDFAFGKTFPGAVMV 243

STANDARD; PRT; 261 AA.

el. 41, Created)
el. 41, Last sequence update)
el. 41, Last annotation update)
factor ligand superfamily member 5 (CD40 ligand) (CD40-
ein).
LG OR CD40L.
tus (Night monkey) (Douroucouli).
aza; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Platyrrhini; Cebidae; Aotinae; Aotus.
5;
N.A.
ytes;
18; PubMed=11491535;
Bostik P., Mayne A.E., King C.L., Genain C.P.,
sari A.A.;
encing, and homology analysis of nonhuman primate
and co-stimulatory molecules.";
53:315-328(2001).
Cytokine that binds to TNFRSF5. Mediates B-cell
ion in the absence of co-stimulus as well as IGE
in the presence of IL-4. Involved in immunoglobulin
ching (By similarity).
omotrimer (By similarity).
R LOCATION: Type II membrane protein. Also exists as an
lar soluble form (By similarity).
cluble form derives from the membrane form by
c processing (By similarity).
: Belongs to the tumor necrosis factor family.

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il to license@isb-sib.ch).

; AAK37542.1; -
; Integral to membrane; ISS.
; F:CD40 receptor binding; ISS.
; P:B-cell proliferation; ISS.
; P:inflammatory response; ISS.
; P:leukocyte cell adhesion; ISS.
; P:platelet activation; ISS.
03263; TNF 5.
06052; TNF family.
08983; TNF like.
03636; TNF_subf.
TNF; 1.
2; CD40LIGAND.
00; TNF 5; 1.
12; TNF_subf; 1.
; TNF; 1.
51; TNF 1; 1.
49; TNF 2; 1.
membrane; Glycoprotein; Signal-anchor.
1 261
MEMBER 5, MEMBRANE FORM.
3 261
TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
MEMBER 5, SOLUBLE FORM (BY SIMILARITY).
1 22
CYTOPLASMIC (POTENTIAL).
3 43
SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
(POTENTIAL).
4 261
EXTRACELLULAR (POTENTIAL).
2 113
CLEAVAGE (BY SIMILARITY).

FT DISULFID 178 218 POTENTIAL.
FT CARBOHYD 240 240 N-LINKED (GLCNAC...) (POTENTIAL).
SQ SEQUENCE 261 AA; 29357 MW; 85E1588B507901B5 CRC64;

Query Match 7.5%; Score 87; DB 1; Length 261;
Best Local Similarity 25.9%; Pred. No. 1.8;
Matches 35; Conservative 21; Mismatches 55; Indels 24; Gs

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Db 136 SVLQWAEKGYTMSNNVLTLENGKQL---TVKRGLYIYIAQVTCSPNEASSQAPF
QY 161 GVALRCLLEF-----SATAASSPQQLRLC-----QVSGLLPLRPGSSLRIRLTF
Db 193 --LCLKPPNRRFERILLRAANTHSSAKP---CGQOSIHLLGGIFELQPGASVFNVTI
QY 210 LKAAPFLTYFGLFQV 224
Db 247 VSHGTGFTSFGLLKL 261

RESULT 12
TNF5 CALJA
ID TNF5 CALJA STANDARD; PRT; 261 AA.
AC Q9BDN3;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Tumor necrosis factor ligand superfamily member 5 (CD40 ligand) (C
L) (CD154 protein).
GN TNFRSF5 OR CD40LG OR CD40L.
OS Callithrix jacchus (Common marmoset).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Callitrichidae;
OC Callithrix.
OX NCBI_TaxID=9483;
RN [1]
SEQUENCE FROM N.A.
RC TISSUE=Lymphocytes;
RX MEDLINE=21383618; PubMed=11491535;
RA Willinger F., Bostik P., Mayne A.E., King C.L., Genain C.P.,
Weiss W.R., Ansari A.A.;
RT "Cloning, sequencing, and homology analysis of nonhuman primate
Fas/Fas-ligand and co-stimulatory molecules.";
RL Immunogenetics 53:315-328(2001).
CC -!- FUNCTION: Cytokine that binds to TNFRSF5. Mediates B-cell
proliferation in the absence of co-stimulus as well as IGE
production in the presence of IL-4. Involved in immunoglobulin
class switching (By similarity).
CC -!- SUBUNIT: Homotrimer (By similarity).
CC -!- SUBCELLULAR LOCATION: Type II membrane protein. Also exists as
extracellular soluble form (By similarity).
CC -!- PTM: The soluble form derives from the membrane form by
proteolytic processing (By similarity).
CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.

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or send an email to license@isb-sib.ch).

CC EMBL; AF344844; AAK37603.1; -
DR HSSP; P29965; IALY.
DR GO; GO:0016021; C:integral to membrane; ISS.
DR GO; GO:0005174; F:CD40 receptor binding; ISS.
DR GO; GO:0042100; P:B-cell proliferation; ISS.
DR GO; GO:0006954; P:inflammatory response; ISS.
DR GO; GO:0007159; P:leukocyte cell adhesion; ISS.
DR GO; GO:0030166; P:platelet activation; ISS.
DR InterPro; IPR003263; TNF_5.

e G.;
utations in X-linked immunodeficiency with hyper-IgM.";
-543(1993).

PRO-155; ASP-211 AND VAL-227.
70; PubMed=7679801;
Mitage R.J., Conley M.E., Rosenblatt H., Jenkins N.A.,
Bedell M.A., Edelhoff S., Distche C.M.,
Faslow W.C., Belmont J.W., Spriggs M.K.;
ene defects responsible for X-linked hyper-IgM
0-993(1993).

ALA-126; ARG-140 AND GLU-144.
38; PubMed=7717401;
la A., Strina D., Sacco M.G., Morali F., Brugnani D.,
ntuano E., Fasth A., Andersson B., Zegers B.J.M.,
znick I., Levy J., Zan-Bar I., Porat Y., Alro P.,
zzoni P., Notarangelo L.D.;
ion of nine novel mutations in the CD40 ligand gene in
X-linked hyper IgM syndrome of various ancestry.";
net. 56:898-906(1995).

PRO-155 AND VAL-227, AND VARIANT ARG-219.
33; PubMed=8550833;
J., Allen R.C., Larche M., Greene J.M., Shigeoka A.O.,
rauf D.C., Belmont J.W., Conley M.E.;
nd conformation polymorphism study of CD40 ligand.
tion analysis and carrier detection for X-linked hyper
t. 97:196-201(1996).

ARG-36; CYS-140; SER-231; MET-254 AND GLY-227 DEL.
77; PubMed=9150729;
himadzu M., Toru H., Seyama K., Nunoi H., Neubauer M.,
h H.D.;
the CD40 ligand gene in 13 Japanese patients with
-IgM syndrome.";
:624-627(1997).

Mediates B-cell proliferation in the absence of co-
s well as IgE production in the presence of IL-4.
n immunoglobulin class switching.

omotrimer.
R LOCATION: Type II membrane protein. Also exists as an
lar soluble form.
CIFICITY: Specifically expressed on activated CD4+
tes.
oluble form derives from the membrane form by
c processing.

ffects in TNF α 5 are the cause of X-linked
ciency with hyper-IgM type 1 (HIGM1) [MIM:308230]. HIGM1
noglobulin isotype switch defect characterized by
oncentrations of serum IgM and decreased amounts of all
ypes. Affected males present at an early age (usually
first year of life) recurrent bacterial and
tic infections, including pneumocystis carinii pneumonia
table diarrhea due to cryptosporidium infection. Despite
on treatment with intravenous immunoglobulin, the
agnosis is rather poor, with a death rate of about 10%
lence.

: Belongs to the tumor necrosis factor family.

NAME=CD40Lbase;

ean CD40L defect database (mutation db);

://www.expasy.org/cd40lbase/";

/ftp.expasy.org/databases/cd40lbase".

NAME=PROW; NOTE=CD guide CD154 entry;

://www.ncbi.nlm.nih.gov/prow/cd/cd154.htm".

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or send an email to license@isb-sib.ch).

CC EMBL; X68550; CAA48554.1; -
CC DR EMBL; 215017; CAA78737.1; -
CC DR EMBL; X67878; CAA48077.1; -
CC DR EMBL; L07414; AAA35662.1; -
CC DR EMBL; D31797; BAA06599.1; -
CC DR EMBL; D31793; BAA06599.1; JOINED.
CC DR EMBL; D31794; BAA06599.1; JOINED.
CC DR EMBL; D31795; BAA06599.1; JOINED.
CC DR EMBL; D31796; BAA06599.1; JOINED.
CC DR PIR; S28017; I53476.
CC DR PDB; 1ALY; 17-SEP-97.
CC DR PDB; 119R; 22-MAY-02.
CC DR Genew; HGNC:11935; TNFSF5.
CC DR MIM; 300386; -
CC DR MIM; 308230; -
CC DR GO; GO:0008887; C:integral to plasma membrane; TAS.
CC DR GO; GO:0005625; C:soluble fraction; TAS.
CC DR GO; GO:0005174; F:CD40 receptor binding; IPI.
CC DR GO; GO:0006916; P:anti-apoptosis; IDA.
CC DR GO; GO:0042100; P:B-cell proliferation; IDA.
CC DR GO; GO:0006954; P:inflammatory response; IDA.
CC DR GO; GO:0045190; P:isotype switching; ISS.
CC DR GO; GO:0007159; P:leukocyte cell adhesion; NAS.
CC DR GO; GO:0030168; P:platelet activation; IDA.
CC DR GO; GO:0007165; P:signal transduction; ISS.
CC DR InterPro; IPR003263; TNF 5.
CC DR InterPro; IPR006052; TNF family.
CC DR InterPro; IPR008983; TNF like.

Query Match 7.4%; Score 86; DB 1; Length 261;
Best Local Similarity 25.9%; Pred. No. 2.2;
Matches 35; Conservative 21; Mismatches 55; Indels 24; G
QY 105 TVSGWEE-----TKINSSSPIDYDRQIGFTVIRAGLYLYCQVHFDGKAVYLKLDI
DB 136 SVLQWAEKGYTMSNNLVLENGKQL---TVKRQGLYTYAQTFCNREASSQAP!
QY 161 GVLAALRCLBEF-----SATAASSPGQLRLC-----QVSGLLPLRPGSSLRITEL
DB 193 --LCLKSPGRFERILLRAANTHSSAKP-----CQQQSIHLGGVFELOFGASVFNVTI
QY 210 LKAAPFLTYFGLFQV 224
DB 247 VSHGTGFTSFGLLKL 261

Search completed: April 7, 2004, 17:45:17
Job time: 10.2829 secs

GenCore version 5.1.6
 yright (c) 1993 - 2004 Compugen Ltd.
 search, using sw model
 7, 2004, 17:41:27 ; Search time 31.3851 Seconds
 (without alignments)
 2261.954 Million cell updates/sec
 1-245-198A-2
 HGLALACGLLVVWSL.....PWAHLKAAPFLTYFGLFQVH 225
 M62
 10.0 , Gapext 0.5
 41 seqs, 315518202 residues
 satisfying chosen parameters: 1017041
 1: 0
 1: 2000000000
 Immun Match 0%
 Immun Match 100%
 ing first 45 summaries

REMBL 25: *
 sp archea: *
 sp bacteria: *
 sp fungi: *
 sp human: *
 sp invertibrate: *
 sp mammal: *
 sp mhc: *
 sp organelle: *
 sp phage: *
 sp plant: *
 sp rodent: *
 sp virus: *
 sp vertebrate: *
 sp unclassified: *
 sp virus: *
 sp bacteriap: *
 sp archesp: *

he number of results predicted by chance to have a
 than or equal to the score of the result being printed,
 by analysis of the total score distribution.

SUMMARIES

Y	h	Length	DB	ID	Description
1	410	11	Q8BXS2	Q8BXS2	Q8BXS2 mus musculus
8	330	4	Q8IZK7	Q8IZK7	Q8IZK7 homo sapien
0	409	5	Q8IGD3	Q8IGD3	Q8IGD3 drosophila
5	409	5	Q8MY88	Q8MY88	Q8MY88 drosophila
3	261	5	Q8MRW2	Q8MRW2	Q8MRW2 drosophila
3	325	5	Q8V5G2	Q8V5G2	Q8V5G2 drosophila
3	415	5	Q8MUJ1	Q8MUJ1	Q8MUJ1 drosophila
9	426	16	Q88IZ6	Q88IZ6	Q88IZ6 pseudomonas
4	684	13	Q7TJ36	Q7TJ36	Q7TJ36 lampetra ja
0	557	16	Q8XQX3	Q8XQX3	Q8XQX3 ralstonia s
8	210	16	Q9A926	Q9A926	Q9A926 caulobacter
8	421	16	Q9HUW2	Q9HUW2	Q9HUW2 pseudomonas
7	213	16	Q82AD2	Q82AD2	Q82AD2 streptomyce
7	1363	5	Q9VFD3	Q9VFD3	Q9VFD3 drosophila
7	287	13	Q90WT9	Q90WT9	Q90WT9 gallus gall
6	224	5	Q9V762	Q9V762	Q9V762 drosophila

17	88	7.6	352	12	089341	089341 hendr
18	88	7.6	353	12	066760	066760 equin
19	88	7.6	532	4	Q16727	Q16727 homo s
20	87.5	7.5	522	10	Q9FTN7	Q9FTN7 oryza
21	87.5	7.5	670	16	Q9AA15	Q9AA15 caulol
22	87	7.5	244	6	Q86227	Q86227 pan tri
23	86.5	7.4	154	16	Q7U7N8	Q7U7N8 syneci
24	86.5	7.4	340	16	Q9HUR8	Q9HUR8 pseud
25	86.5	7.4	504	16	Q92KA4	Q92KA4 rhizol
26	86.5	7.4	1237	11	Q7TPS4	Q7TPS4 mus m
27	86	7.4	260	10	Q8S2N9	Q8S2N9 oryza
28	85.5	7.4	331	10	Q942P9	Q942P9 oryza
29	85.5	7.4	724	5	Q868S9	Q868S9 anophe
30	85.5	7.4	2841	10	Q7XU06	Q7XU06 oryza
31	85	7.3	116	16	Q7V2A2	Q7V2A2 proch
32	84.5	7.3	377	16	Q7UEA6	Q7UEA6 rhodo
33	84.5	7.3	422	16	Q9RKB0	Q9RKB0 strep
34	84.5	7.3	430	2	Q9REU1	Q9REU1 strept
35	84.5	7.3	1079	13	Q8UVR4	Q8UVR4 xenop
36	84.5	7.3	1118	16	Q98E34	Q98E34 rhizo
37	84	7.2	2962	5	Q93326	Q93326 caenor
38	83.5	7.2	394	16	Q92V66	Q92V66 rhizo
39	83	7.1	467	16	Q9S2Y4	Q9S2Y4 strep
40	82.5	7.1	174	16	Q9CKX1	Q9CKX1 paste
41	82.5	7.1	314	5	Q8WPH7	Q8WPH7 theile
42	82.5	7.1	501	16	Q89IN3	Q89IN3 brady
43	82.5	7.1	549	16	Q8RC38	Q8RC38 therm
44	82.5	7.1	718	6	Q8HXH0	Q8HXH0 macaca
45	82	7.1	619	5	Q8SUH9	Q8SUH9 enceph

ALIGNMENTS

RESULT 1
 Q8BXS2 PRELIMINARY; PRT; 410 AA.
 ID Q8BXS2
 AC Q8BXS2;
 DT 01-MAR-2003 (TREMBLrel. 23, Created)
 DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
 DE Tumor necrosis factor.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Retina;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium,
 RA the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation
 RT 60,770 full-length cDNAs";
 RL Nature 420:563-573(2002).
 DR ENBL; AK044387; BAC31897.1; -.
 DR PIR; PT0714; PT0714.
 DR GO; GO:0016020; C:membrane; IEA.
 DR GO; GO:0005164; F:tumor necrosis factor receptor binding; IEA.
 DR GO; GO:0006955; P:immune response; IEA.
 DR InterPro; IPR006052; TNF_family.
 DR SMART; SM00207; TNF; 2.
 DR PROSITE; PS00251; TNF_1; 1.
 DR PROSITE; PS00049; TNF_2; 2.
 SQ SEQUENCE 410 AA; 45681 MW; 590A4B74C33FB8D4 CRC64;
 Query Match 82.1%; Score 954.5; DB 11; Length 410;
 Best Local Similarity 88.6%; Pred. No. 1.3e-83;
 Matches 195; Conservative 1; Mismatches 13; Indels 11; Gap
 QY 1 VLSIGLALCLGLLVVWSLGSWATLSA-QEPSEELTAEDRRPELNPOTERSQD

GLALACGLLLVSVLSGWSATLSAQEPSELTAEADRRPELNPQTESQDVVP 84
 LVPRPSAPKGRKAPRAIAAHYVHPRPGDGAQAGVDGTSGWEETKINSASP 119
 LVPRPSAPKGRKAPRAIAAHYVHPRPGDGAQAGVDGTSGWEETKINSASP 144
 RQIGFTVIRAGLYLYCQVHFDGKAVYLKLDLLVNGVLAALRCLEEFSAATAASP 179
 RQIGFTVIRAGLYLYCQVHFDGKAVYLKLDLLVNGVLAALRCLEEFSAATAASP 204
 RLCQVSGLLPLR-----PGSSLRITLPAWHL 210
 RLCQTE-LQSURREVSRLQRSGPQKQGERPQSL 243

RELIMINARY; PRT; 330 AA.

TrEMBLrel. 23, Created
 TrEMBLrel. 23, Last sequence update
 TrEMBLrel. 25, Last annotation update
 (Human).
 azoia; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Primates; Carnivora; Hominidae; Homo.
 6;

N.A.
 24; PubMed12411489;
 B., Medina J.P., Lopez-Fraga M., Lozano J.C.,
 M., Picard A., Martinez-A C., Garcia-Sanz J.A.,
 hybrid mRNA encodes TWE-PRIL, a functional cell surface
 protein.
 1-5720(2002).
 ; AAL50443.1;
 ; C-membrane; IEA.
 ; F-tumor necrosis factor receptor binding; IEA.
 ; P-immune response; IEA.
 06052; TNF family.
 08983; TNF-like.
 TNF; 1.
 ; TNF; 1.
 51; TNF 1; 1.
 49; TNF 2; 2.
 AA; 36588 MW; FC6F3BCA29C029AE CRC64;
 rity 52.8%; Score 613; DB 4; Length 330;
 nservative 8; Mismatches 14; Indels 0; Gaps 0;

ALACGLLLVSVLSGWSATLSAQEPSELTAEADRRPELNPQTESQDVVPFL 61
 ALACGLLLVSVLSGWSATLSAQEPSELTAEADRRPELNPQTESQDVVPFL 85
 PRPSAPKGRKAPRAIAAHYVHPRPGDGAQAGVDGTSGWEETKINSASP 121
 PRPSAPKGRKAPRAIAAHYVHPRPGDGAQAGVDGTSGWEETKINSASP 145
 3EFTVIRAGLYLYCQ 142
 3EFTVIRAGLYLYCQ 166

RELIMINARY; PRT; 409 AA.

EMBLrel. 23, Created
 EMBLrel. 23, Last sequence update
 EMBLrel. 25, Last annotation update

BCDNA:RH51659.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Y;
 RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
 RA Champe M., Chavez C., Dorsett V., Dresnek D., Farfan D., Frise E.,
 RA George R., Gonzalez M., Guarin H., Krommiller B., Li P., Liao G.,
 RA Miranda A., Mungall C.J., Nunoo J., Pacleb J., Paragas V., Park S.,
 RA Patel S., Phouanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.
 RA Celniker S.;
 RL Submitted (NOV-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; BT001838; AAN71595.1;
 DR FlyBase; FBgn0064801; BCDNA:RH51659.
 DR GO; GO:0016020; C-membrane; IEA.
 DR GO; GO:0005164; F-tumor necrosis factor receptor binding; IEA.
 DR GO; GO:0006955; P-immune response; IEA.
 DR InterPro; IPR006052; TNF family.
 DR InterPro; IPR008983; TNF-like.
 DR SMART; SM00207; TNF; 1.
 DR PROSITE; PS00251; TNF 1; 1.
 DR PROSITE; PS0049; TNF 2; 1.
 SQ SEQUENCE 409 AA; 46401 MW; FC2E9BD9E012D257 CRC64;

Query Match 10.0%; Score 116.5; DB 5; Length 409;
 Best Local Similarity 24.3%; Pred. No. 0.01;
 Matches 50; Conservative 32; Mismatches 87; Indels 37; G;
 QY 29 QBPSPQELTAEDRRPELNPQTESQDVVPFLQVLRPRRSPAPKGRAPRAIA/
 Db 231 QEKSSNEATSKERPAHLHRRHGRH-----RHLLVRKARS-----EDSRP-----AJ
 QY 89 VHRPFGDGAQAGVDGTSGWEETKINSPLRDRGCEFTVIRAGLYLYCQVH
 Db 278 LSSRRRHQGSN-GYHGDVYIGNDNERNVYQ-HFQTRDGLVTWTGLYVVAQIC
 QY 149 KAVYLKLDLLVNGVLA-----LRCLEEFSAATAASPPQLRLQVSGLLPLRPG
 Db 336 HD-----QNGFTVFGQDTPFLQCLN---TVPTNMPKHVHTCHTSLIHLERNE
 QY 202 IRTL---PWAHLKAPFLTYGLFOV 224
 Db 384 LKDIHNDENAVLRGNRNSYFGIFKV 409

RESULT 4
 Q8MY88 PRELIMINARY; PRT; 409 AA.
 ID Q8MY88
 AC Q8MY88;
 DT 01-OCT-2002 (TrEMBLrel. 22, Created)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE TNF superfamily ligand, Eiger (Tumor necrosis factor family member
 DE DT1).
 DE GN
 GN EIGER OR DT1 OR CG:2919.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=22060500; PubMed=12065414;
 RA Igaki T., Kanda H., Yamamoto-Goto Y., Kanuka H., Kuranaga E.,
 RA Aigaki T., Miura M.;
 RT "Eiger, a TNF superfamily ligand that triggers the Drosophila JNK
 RT pathway".
 RL ENBO J. 21:3009-3018(2002).
 RN [2]

Query Match	9.3%	Score 107.5	DB 5	Length 261
Best Local Similarity	23.1%	Pred. No. 0.042		
Matches	48	Conservative	34	Mismatches 91; Indels 35; Gaps 1
QY	29	QEPQOEELTAEDRRPELPINPQTEESQDVVFLEQVLVRRPS--APGKRKARPRRAIA		
Db	77	QEKSNSEATSKESAPLHRRRMSRH-----RHLLVRKGESLLISARSDSRP-----A		
QY	87	YEVHPRPQDGAQAGVDGTVSGWEEETKINSSPLRYDROIGETVIRAGLYLYLYCQVH		
Db	128	PHLSRRRHQSGM--GYGDMYIGNDNERNYQG-HFQTRDGLVITNTGLIYYVYQALIC		
QY	147	EKGAVYMLDLVNVGLA-----LRLCEFSATAASFGPQLRQCVSGLLPLRPG		
Db	186	NSHD-----QNGFIVQGDTPFLQCLN---TVPTNPHKVHTCHTSLGLIHLENR		
QY	200	LRIRTL-----PWAHLKAAFPFLTYGFLQV 224		
Db	234	HLKDIHNDRNAVLREGNRRSYGIFKV 261		
RESULT 6				
Q9V5G2	PRELIMINARY:	PRT:	325	AA.
ID	Q9V5G2			
AC	Q9V5G2			
CD	01-MAY-2000 (TrEMBLrel. 13, Created)			
DT	01-MAY-2000 (TrEMBLrel. 13, Last sequence update)			
DT	01-OCT-2003 (TrEMBLrel. 25, Last annotation update)			
DE	CG12919 protein.			
GN	EIGER OR CG12919.			
OS	Drosophila melanogaster (Fruit fly).			
OC	Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;			
OC	Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;			
OC	Ephydroidea; Drosophilidae; Drosophila.			
OX	NCBI_TaxID=7227;			
RN	[1]			
RC	SEQUENCE FROM N.A.			
RP	STRAIN=Berkeley;			
RX	MEDLINE=20196006; PubMed=10731132;			
RA	Adams M.D., Cainick S.E., Holt R.A., Evans C.A., Gocayne J.D.,			
RA	Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,			
RA	George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,			
RA	Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,			
RA	Brandon R.C., Rogers J.-H.C., Blazej R.G., Champagne M., Pfeiffer B.D.,			
RA	Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.			
RA	Abrial J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.			
RA	Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley B.M.,			
RA	Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,			
RA	Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,			
RA	Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra J.			
RA	Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,			
RA	de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,			
RA	Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn			
RA	Durbin K.J., Evangelista C.C., Ferraz C., Ferriera S., Fleischmann			
RA	Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,			
RA	Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,			
RA	Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,			
RA	Hosdin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,			
RA	Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.			
RA	Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,			
RA	Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,			
RA	Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,			
RA	Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,			
RA	Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,			
RA	Nelson D.R., Nelson K.A., Nixon K., Nussekern D.R., Pacleb J.M.,			
RA	Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.			
RA	Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,			
RA	Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,			
RA	Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,			
RA	Swirkas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,			
RA	Wang Z.-Y., Wassarman D.A., Weinstock G.M., Weissbach J.,			
RA	Williams S.M., Woodgate T., Worley K.C., Wu D., Yang S., Yao Q.A.,			
RA	Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.			

hong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 yers E.W., Rubin G.M., Venter J.C.;
 sequence of *Drosophila melanogaster*.
 185-2195(2000).
 1; AAF58848.1; -.
 0033483; eiger.
 0; C-membrane; IEA.
 4; F-tumor necrosis factor receptor binding; IEA.
 06052; TNF family.
 008983; TNF-like.
 7; TNF; 1.
 251; TNF 1; 1.
 049; TNF 2; 1.
 5 AA; 36862 MW; 685CCB69694FIA3A CRC64;
 9.3%; Score 107.5; DB 5; Length 325;
 arity 23.1%; Pred. No. 0.056;
 conservative 34; Mismatches 91; Indels 35; Gaps 9;
 QEELTAEDRREPPPELNPQTESQDVVFLQLVPRRS--APGKRPRAIAAH 86
 SNEATSKESAPLHRRHRSR-----RHLLVRKGSLLSARSDSRP----AAH 191
 PRPQDGAQAGVGTSGWETKINSPLRYDRQIGFTVIRAGLYLYCYVHD 146
 SRRHQGSM-GYHGDVMIYGNDRNSYQG-HFQTRDGLVLTNTGLVYVYAAICYN 249
 VYLKDLLVNGVLA-----LRCLFEFSATAASSPGQLRLCQVGLPLRPGSS 199
 -----QNGFTVFGQDTFFLQCLN-----TVPTNMPKHVHTCHTSLIHLNER 297
 TL---PWAHLKAAPFLTYFGLFOV 224
 IHNDRNAVLRGNRRSYFGIFKV 325
 PRELIMINARY; PRT; 415 AA.
 REMBLrel. 22, Created)
 REMBLrel. 22, Last sequence update)
 REMBLrel. 25, Last annotation update)
 19 OR DARTH.
 anogaster (fruit fly).
 azoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 pterygota; Diptera; Brachycera; Muscomorpha;
 rosophilidae; Drosophila.
 7;
 N.A.
 23; PubMed=12176339;
 M., Baer K.;
 TNF Signaling Mechanisms. JNK-Dependent Apoptosis
 iger, the *Drosophila* Homolog of the TNF Superfamily.";
 :1263-1268(2002).
 N.A.
 38; PubMed=12894227;
 aaty W.S., Chen P., Tomar R.S., Eby M.T., Chappo J.,
 re N., Zachariah S., Sinha S.K., Abrams J.M.,
 ; receptor, Wengen, comprise a TNF-like system in
 60-4867(2003).
 ; AAM76710.1; -.
 ; AAM66763.1; -.
 033483; eiger.
 ; C-membrane; IEA.
 ; F-tumor necrosis factor receptor binding; IEA.
 ; P-immune response; IEA.

DR InterPro: IPR006052; TNF family.
 DR InterPro: IPR008983; TNF-like.
 DR SMART: SM00207; TNF_1.
 DR PROSITE: PS00251; TNF_1; 1.
 DR PROSITE: PS0049; TNF_2; 1.
 SQ SEQUENCE 415 AA; 46918 MW; E087A26DE222DBF CRC64;
 Query Match 9.3%; Score 107.5; DB 5; Length 415;
 Best Local Similarity 23.1%; Pred. No. 0.077;
 Matches 48; Conservative 34; Mismatches 91; Indels 35; G
 QY 29 QEPSQEELTAEDRREPPPELNPQTESQDVVFLQLVPRRS--APGKRPRAIAAH 86
 Db 231 QEKSSNEATSKESAPLHRRHRSR-----RHLLVRKGSLLSARSDSRP---- 191
 QY 87 YEVHPRPQDGAQAGVGTSGWETKINSPLRYDRQIGFTVIRAGLYLYCYVHD 146
 Db 282 FHLSRRHQGSM-GYHGDVMIYGNDRNSYQG-HFQTRDGLVLTNTGLVYVYAAICYN 249
 QY 147 EGKAVYLKDLLVNGVLA-----LRCLFEFSATAASSPGQLRLCQVGLPLR 199
 Db 340 NSHD-----QNGFTVFGQDTFFLQCLN-----TVPTNMPKHVHTCHTSLIHL 297
 QY 200 LRIRTL---PWAHLKAAPFLTYFGLFOV 224
 Db 388 IHLKDIHNDRNAVLRGNRRSYFGIFKV 325
 PRELIMINARY; PRT; 426 AA.
 Q88126
 ID Q88126 PRELIMINARY; PRT; 426 AA.
 AC Q88126;
 DT 01-JUN-2003 (T-EMBLrel. 24, Created)
 DT 01-JUN-2003 (T-EMBLrel. 24, Last sequence update)
 DT 01-JUN-2003 (T-EMBLrel. 24, Last annotation update)
 DE Conserved hypothetical protein.
 GN PP2853.
 OS Pseudomonas putida (strain KT2440).
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 OC Pseudomonadaceae; Pseudomonas.
 OX NCBI_TaxID=160488;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=22423060; PubMed=12534463;
 RA Nelson K.E., Weinel C., Paulsen I.T., Dodson R.J., Hilbert H.,
 RA Martins dos Santos V.A.P., Fouts D.E., Gill S.R., Pop M., Holmes I.
 RA Brinkac L., Beanan M., DeBoy R.T., Daugherty S., Kolonay J.,
 RA Madupu R., Nelson W., White O., Peterson J., Khouri H., Hance I.,
 RA Chris Lee P., Holtzapple E., Scanlan D., Tran K., Moazzez A.,
 RA Utterback T., Rizzo M., Lee K., Kosack D., Moestl D., Wedler H.,
 RA Lauber J., Stjepandic D., Hoheisel J., Straetz M., Heim S.,
 RA Kiewitz C., Eisen J., Timmis K.N., Duisterhoft A., Thummel B.,
 RA Fraser C.M.;
 RT "Complete genome sequence and comparative analysis of the
 RT metabolically versatile *Pseudomonas putida* KT2440.";
 RL Environ. Microbiol. 4:799-808(2002).
 DR EMBL; AF016784; AAN68461.1; -.
 DR TIGR; PP2853; -.
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 426 AA; 46020 MW; FEDC7E266C982633 CRC64;
 Query Match 8.9%; Score 103; DB 16; Length 426;
 Best Local Similarity 29.1%; Pred. No. 0.22;
 Matches 50; Conservative 17; Mismatches 65; Indels 40; G
 QY 7 ALACLGILLVVSLGSAQPSQELTAEDRREPPPELNPQTESQDVVFLQL 86
 Db 9 AFVCLTTTLAPATAL-----YAADPQVEAL-----RQELIELKRRYEAQQALMWLE 146
 QY 67 PRRSAPGKRPRAIAAHVEVHPRPQDGAQ-----AGVDGTV-SGWETKINS 191
 Db 60 QVEAPAA--AQPKRLVKS-----PAGVKGAQTVASGAPGTGGTSSYGOALTADSE 249

q

9


```
QY      135 GUYIYCQHFD-----EGKAVYLKDILVNGVUALRCLBEFS
          |||:|||
Db       185 GKYYVYSQIFYFRYSDGAGRVSPQLVCINWKTYSQPILLIKGV
          |||:|||
QY      176 ASSPQG--LRLCQVGSLLPDPGSSLIRTLPAWLKAAPFLYFGLFQV 224
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06:25:13 2004

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April 7, 2004, 17:46:43
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16:25:12 2004

us-09-245-198a-2.rag

E

GenCore version 5.1.6
copyright (c) 1993 - 2004 CompuGen Ltd.

n search, using sw model

il 7, 2004, 17:37:32 ; Search time 45.0884 Seconds
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09-245-198A-2

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SUM62

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6107 segs, 282547505 residues

s satisfying chosen parameters: 1586107

th: 0
th: 2000000000

nimum Match 0%
ximum Match 100%
string first 45 summaries

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Genesecp1980s: *
Genesecp1990s: *
Genesecp2000s: *
Genesecp2001s: *
Genesecp2002s: *
Genesecp2003as: *
Genesecp2003bs: *
Genesecp2004s: *

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
id by analysis of the total score distribution.

SUMMARIES

Seq	Length	DB	ID	Description
1.0	225	2	AAW47524	AAW47524 Mus muscu
1.0	225	3	AB07527	AB07527 Amino aci
1.0	249	7	ADC97712	ADC97712 Murine FL
1.7	211	2	AAW93591	AAW93591 Mouse TNF
1.8	249	2	AAW29745	AAW29745 TNF relat
1.8	249	2	AAU09369	AAU09369 Human tum
1.8	249	3	AAU95338	AAU95338 Human PRO
1.8	249	3	AB07526	AB07526 Amino aci
1.8	249	4	AAE00891	AAE00891 Human TRE
1.8	249	5	AAU86129	AAU86129 Human PRO
1.8	249	6	ABR42315	ABR42315 Human TRE
1.8	249	7	ADC35206	ADC35206 Human TNF
1.8	284	2	AAW47525	AAW47525 Homo sapi
1.8	208	2	AAW93590	AAW93590 Human TNF
1.8	273	4	AAU03499	AAU03499 TWEAK ext
1.9	189	2	AAW29746	AAW29746 TNF relat
1.9	189	4	AAE00892	AAE00892 Human UL4
2.0	146	4	AAE00895	AAE00895 Human TRE
3.6	406	5	AAU77717	AAU77717 Drosophil
3.5	409	5	AAU77718	AAU77718 Drosophil
3.3	325	4	ABW67553	ABW67553 NF-kB rec
3.2	294	2	AAW69956	AAW69956 NF-kB rec
3.2	294	2	AAW68292	AAW68292 NF-kB rec
3.2	294	2	AAE08737	AAE08737 Murine re
3.2	294	4	AAE04425	AAE04425 Murine re

26	106.5	9.2	294	4	AAE01992	Mu
27	106.5	9.2	294	5	AAE26102	Mol
28	106.5	9.2	294	7	ADB16986	Mu
29	106.5	9.2	294	7	ADC73000	Mu
30	106.5	9.2	294	7	ADC78266	Mur
31	103	8.9	220	4	AAE62340	GpJ
32	103	8.9	426	6	ABU39962	PrC
33	102.5	8.8	316	2	AAW83017	Ost
34	102.5	8.8	316	2	AAW83194	Hun
35	102.5	8.8	316	2	AAW59654	Ami
36	102.5	8.8	316	2	AAU17874	Mur
37	102.5	8.8	316	3	AAU91024	Mol
38	102.5	8.8	316	3	AAU84418	Ami
39	102.5	8.8	316	3	AAU84419	Ami
40	102.5	8.8	316	5	AAU78289	Mol
41	102.5	8.8	316	6	ABR42071	Hun
42	102.5	8.8	316	6	ABW99477	Ami
43	102.5	8.8	316	6	ABU08463	Ami
44	102.5	8.8	316	6	ABR55560	Ami
45	99	8.5	234	4	AAE62339	GpJ

ALIGNMENTS

RESULT 1
AAW47524
ID AAW47524 standard; protein; 225 AA.
XX AC AAW47524;
XX AC
XX 21-JUL-1998 (first entry)
XX DE Mus musculus tumour necrosis factor related ligand (TRELL).
XX DE TRELL; tumour necrosis factor related ligand; tnfr; treatment; cancer
XX KW autoimmune disease; immune system; stimulation; suppression;
XX KW graft rejection.
XX OS Mus musculus.
XX FH Key Location/Qualifiers
FT Domain 1..21
FT /note= "hydrophobic, transmembrane domain"
XX PN WO9805783-A1.
XX PD 12-FEB-1998.
XX PF 07-AUG-1997; 57WO-US013945.
XX PR 07-AUG-1996; 96US-0023541P.
XX PR 18-OCT-1996; 96US-0028515P.
XX PR 18-MAR-1997; 97US-0040820P.
XX PA (BIOJ) BIOGEN INC.
XX PA (UYGE-) UNIV GENEVA FACULTY MEDICINE.
XX PI Chicheportiche Y, Browning JL;
XX DR WPI; 1998-145619/13.
XX DR N-PSDB; AAV18599.
XX PT Tumour necrosis factor related ligand - useful for, e.g. treating
XX PT auto-immune disease and immune responses to tissue grafts.
XX PS Claim 12; Page 48-50; 69pp; English.
XX CC The sequence is that of mouse tumour necrosis factor related ligand
XX CC (TRELL). TRELL or active fragments can be included with a carrier
XX CC pharmaceutical compositions to treat cancer, autoimmune diseases,
XX CC immune responses to tissue grafts, or to stimulate or suppress the
XX CC system. It is useful to screen for TRELL receptors, by labelling

bel and screening compositions for binding. Agents with TRELL-receptor binding can also be screened for, can be screened, optionally with interferon- γ , to induce cell death, suppress or alter immune responses (especially involving T cells, B cells, or natural killer cells) in the presence of T cells (e.g., T cells) or B cells (e.g., B cells) or natural killer cells (e.g., natural killer cells). It's coding sequence can be used in gene therapy for TRELL-related diseases in mammals (especially humans), e.g., tumours, and inflammatory diseases or inherited genetic disorders, by introducing the sequence into cells, and expressing, therapeutically effective amounts of the sequence in cells, e.g., a virus comprising a gene encoding TRELL. It may also be used in the preparation of probe probes for screening genetic DNAs for TRELL-encoding sequences and for antisense

AA;
 Query Match 100.0%; Score 1162; DB 2; Length 225;
 Best Local Similarity 100.0%; Pred. No. 3.1e-112;
 Matches 225; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 1 VLISGLALACIGLLVSVLSGWSATLSAQPSQELTAEDRREPPELNPTQESQI
 1 VLISGLALACIGLLVSVLSGWSATLSAQPSQELTAEDRREPPELNPTQESQI
 61 LEQVRRPSAPKGRKARPRRAIAAHVEVHPRGQDGAQAGVDTVSGWEETKINE
 61 LEQVRRPSAPKGRKARPRRAIAAHVEVHPRGQDGAQAGVDTVSGWEETKINE
 121 RYDQIGFTVIRAGLYLYCOVHFDEGKAVYKLDLLVNGVLALRCLEEPSATAJ
 121 RYDQIGFTVIRAGLYLYCOVHFDEGKAVYKLDLLVNGVLALRCLEEPSATAJ
 181 POLRLCQVSGLLPLRPGSSLRITLPAHLKAAPFLTYFGLFQVH 225
 181 POLRLCQVSGLLPLRPGSSLRITLPAHLKAAPFLTYFGLFQVH 225
 LCQVSGLLPLRPGSSLRITLPAHLKAAPFLTYFGLFQVH 225
 LCQVSGLLPLRPGSSLRITLPAHLKAAPFLTYFGLFQVH 225

Standard; protein; 225 AA.

(first entry)

Sequence of a soluble recombinant murine TWEAK protein.

Immunological disorder; immune response; inflammation; agent; autoimmune disease; organ transplant rejection; Host disease; GVHD; lymphoid cell malignancy; shock; tumour.

1.

2000WO-US001044.

99US-0116168P.

EN INC.

5036/41.

and treating immune responses using modulators, especially of TWEAK, TWEAK receptors and TWEAK ligands, useful for inflammation and graft versus host disease.

Fig 1; 45pp; English.

The present sequence represents a TWEAK protein. The specific method describes a method for preventing or treating an immunological disorder by administering a TWEAK blocking agent. The method may be used for preventing and/or activity of TWEAK. These disorders include autoimmune diseases, acute and chronic inflammation, organ transplant rejection, Graft-versus-Host disease (GVHD), lymphoid cell malignancies, sepsis, other forms of shock, loss of immune responsiveness (as seen in immunodeficiency virus (HIV) infections) and failure of the immune response to tumour growth

Sequence 225 AA;

Query Match 100.0%; Score 1162; DB 3; Length 225;
 Best Local Similarity 100.0%; Pred. No. 3.1e-112;
 Matches 225; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 1 VLISGLALACIGLLVSVLSGWSATLSAQPSQELTAEDRREPPELNPTQESQI
 1 VLISGLALACIGLLVSVLSGWSATLSAQPSQELTAEDRREPPELNPTQESQI
 61 LEQVRRPSAPKGRKARPRRAIAAHVEVHPRGQDGAQAGVDTVSGWEETKINE
 61 LEQVRRPSAPKGRKARPRRAIAAHVEVHPRGQDGAQAGVDTVSGWEETKINE
 121 RYDQIGFTVIRAGLYLYCOVHFDEGKAVYKLDLLVNGVLALRCLEEPSATAJ
 121 RYDQIGFTVIRAGLYLYCOVHFDEGKAVYKLDLLVNGVLALRCLEEPSATAJ
 181 POLRLCQVSGLLPLRPGSSLRITLPAHLKAAPFLTYFGLFQVH 225
 181 POLRLCQVSGLLPLRPGSSLRITLPAHLKAAPFLTYFGLFQVH 225

RESULT 3

ADC97712

ID ADC97712 standard; protein; 249 AA.

XX AC ADC97712;

XX DT 15-JAN-2004 (first entry)

XX DE Murine FL-TWEAK.

XX KW Murine; FL-TWEAK; TNF relatedness and weak ability to induce cell death; Tumour Necrosis Factor; TWEAK; fibrosis; cardiac disease; liver disease; lung disease; kidney disease; skin disease; skeletal muscle disease; adipose tissue disease; gastrointestinal tract disease; pancreatic disease; reproductive organ disease; neural disease; cartilage disease; bone disease; connective tissue disease; cellular death; hepatocellular carcinoma; dermatological; gastrointestinal; osteopathic.

XX OS Mus sp.

XX PN WO2003086311-A2.

XX PD 23-OCT-2003.

XX PF 09-APR-2003; 2003WO-US011350.

XX PR 09-APR-2002; 2002US-0371611P.

XX PA (BIOJ) BIOGEN INC.

XX PI Burkly L, Jakubowski A, Zheng T, Hahn K;

XX DR WPI; 2003-845256/78.

XX DR N-PSDB; ADC97713.

XX PT Treating a TWEAK-related condition, e.g. liver, gastrointestinal bone, pancreatic, cartilage or neural tissue condition in a subject.

nistering to the subject a TWEAK agonist or antagonist.

ID NO 1; 120pp; English.

quence is murine transmembrane FL-TWEAK (TNF relatedness
ty to induce cell death, where TNF is Tumour Necrosis
is a member of the TNF family. TWEAK agonists or
e useful for treating a TWEAK-related condition, e.g.
iac disease; liver disease; lung disease; kidney disease;
skeletal muscle disease; adipose tissue disease;
al tract disease; pancreatic disease; reproductive organ
l disease; cartilage disease; bone disease; connective
; cellular death; and a pathological condition of a tissue
WEAK receptor.

A;

100.0%; Score 1162; DB 7; Length 249;
rity 100.0%; Pred. No. 3.6e-112;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

LALACLGLLLVSVLSGWTLSAQEPSOEELTAEDRPELNPQTEESQDVVPF 60
LALACLGLLLVSVLSGWTLSAQEPSOEELTAEDRPELNPQTEESQDVVPF 84

RPRRSPKGRKARPRRAIAAHVEVHPRGDGAQAGVDGTGSGWEETKINSSPL 120
RPRRSPKGRKARPRRAIAAHVEVHPRGDGAQAGVDGTGSGWEETKINSSPL 144

IGFTVIRAGLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLBEFSATASSPG 180
IGFTVIRAGLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLBEFSATASSPG 204

COVSGLLPLRGSSLRIRTLPAHLKAAPFLTYFGLFQVH 225
COVSGLLPLRGSSLRIRTLPAHLKAAPFLTYFGLFQVH 249

lard; protein; 211 AA.

(first entry)

otein.

s factor receptor; signal transducer molecule; TNF; APO4;
abnormality; gestational abnormality; prostate cancer;
09; TNRL-1; TNRL-3; diagnosis; treatment; therapy; disease;
main; immunogen; antibody preparation; breast carcinoma;
ise.

98WO-US018393.

97US-00924634.

WASHINGTON.

191/17.

425.

rosis Factor family receptor polypeptides and ligands -
agnosis and treatment of prostate cancer and developmental

or gestational abnormalities.

Claim 40; Fig 13B; 156pp; English.

This invention describes isolated Tumor Necrosis Factor (TNF) fami
receptor polypeptides: APO4, APO6, APO8 and APO9 or their active
fragments, and isolated TNF related ligands 1 and 3 (TNRL1 and TNF
their active fragments. APO4 is useful for diagnosing prostate car
determining levels of APO4 in an individual. Prostate cancer can
treated using APO4 selective binding agents linked to a therapeutic
moisty. APO4 polypeptides are also useful for identifying selectiv
binding agents, useful in diagnosis/treatment of disease by bindi
agents to the polypeptide/active fragment which is extracellular,
expressed on the cell surface. The binding is preferably performe
vivo. APO4 polypeptides/ active fragments are also useful for scr
for agonists and antagonists by binding and observing the changer
activity. Effective pharmacological agents useful in diagnosis or
treatment of disease are also identified using APO4 polypeptides/
fragments and APO4 signal transducer molecules that specifically
with a cytoplasmic domain of APO4 and detecting a change in level
activity. The method is performed in vivo or in vitro. APO polype
are all useful as immunogens for preparing antibodies. APO4 is al
useful for diagnosis/treatment of developmental or gestational
abnormalities. APO8 was transfected to human breast carcinoma cel
MCF-7, and induced apoptosis

Sequence 211 AA;

Query Match 93.7%; Score 1089; DB 2; Length 211;
Best Local Similarity 99.5%; Pred. No. 1.1e-104;
Matches 210; Conservative 0; Mismatches 1; Indels 0; G

QY 15 LVVSLGSGWTLSAQEPSOEELTAEDRPELNPQTEESQDVVPFLEQLVPRRS;

Db 1 LVVSLGSGWTLSAQEPSOEELTAEDRPELNPQTEESQDVVPFLEQLVPRRS;

QY 75 RKARPRRAIAAHVEVHPRGDGAQAGVDGTGSGWEETKINSSPLRYDQIGFT

Db 61 RKARPRRAIAAHVEVHPRGDGAQAGVDGTGSGWEETKINSSPLRYDQIGFT

QY 135 GLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLBEFSATASSPGRLCQVSG

Db 121 GLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLBEFSATASSPGRLCQVSG

QY 195 RPSGSLRIRTLPAHLKAAPFLTYFGLFQVH 225

Db 181 RPSGSLRIRTLPAHLKAAPFLTYFGLFQVH 211

RESULT 5

AAW29745

ID AAW29745 standard; protein; 249 AA.

XX AC AAW29745;

XX DT 27-OCT-1998 (first entry)

XX DE TNF related endothelium proliferative agent protein.

XX KW TNF; endothelium proliferative agent; TREPA; wound healing; cancer
tissue grafting; vascularisation; apoptosis; autoimmune; birth co

XX OS Homo sapiens.

XX PN WO9835061-A2.

XX PD 13-AUG-1998.

XX PF 12-FEB-1998; 98WO-US002859.

XX PR 12-FEB-1997; 97US-00798692.

XX PR 10-FEB-1998; 98US-00021706.

XX

P LAB.

255/38.

613.

leic acid encoding TREPA - useful for diagnosis and autoimmune disease, tumours and inflammation.

e 123-4; 142pp; English.

ed endothelium proliferative agent (TREPA), or its agonists, are used to treat a deficit of TREPA, e.g. to healing or tissue grafting, by promoting vascularisation, e apoptosis for treating cancer and eliminating autoreactive n adjunct to cancer chemotherapy or antiviral treatment. s can also be used to target cytotoxic agents or for ation of the corresponding receptor, the nucleic acid for used to transform tumour cells to render them more. TREPA and to screen for TREPA mimics. Ribozymes, antisense ies or peptides, are used to treat TREPA-associated . tumours and metastases (by inhibiting vascularisation), or a wide range of autoimmune conditions, conditions ormal stimulation of epithelial cells (e.g. is), for birth control (inhibiting ovulation and placental other angiogenic conditions (e.g. ulcers)

AA;

arity 87.8%; Score 1020; DB 2; Length 249;

conservative 9; Mismatches 16; Indels 0; Gaps 0;

LALACLGLLVVSLGSWATLSAQPSQBELTAEDRRPELNPQTEESQDVPEFL 61

LALACLGLLVVSLGSRASLSAQEPQAQELVAEDQPSLNPQTEESQDPAPFL 85

RPRSPAPKGRKPRRAIAAHYVHPRPQDGAQAGVDGTVSGWEEKINSPLR 121

RPRSPAPKGRKTRARRAIAAHYVHPRPQDGAQAGVDGTVSGWEEKINSPLR 145

IGFTVIRAGLYLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLEEFSAASP 181

IGFEFTVIRAGLYLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLEEFSAASP 205

QVSGLLPLRPGSSLRIRITLPWAHLKAAPFLTYFGLFQVH 225

QVSGLLALRPGSSLRIRITLPWAHLKAAPFLTYFGLFQVH 249

dard; protein; 249 AA.

(first entry)

necrosis factor Apo-3 ligand protein sequence.

; necrosis factor; Apo-3 ligand; lymphotoxin; apoptosis; dependent transcription; JNK/SAPK-dependent response; cancer.

98WO-US021407.

97US-0062037P.

PR 17-DEC-1997; 97US-0069862P.
XX (GETH) GENENTECH INC.
PA Ashkenazi AJ, Marsters SA, Pitti R;
XX WPI; 1999-287982/24.
XX DR N-PSDB; AAX56000.
XX PT New human Apo3- ligand (a tumor necrosis factor) homologue.
XX Claim 1; Fig 1; 74pp; English.
XX The present sequence represents a human tumour necrosis factor (T lymphotoxin homologue designated Apo-3 ligand. Apo-3 ligand has cytostatic activity. Apo-3 ligand can be used to induce apoptosis mammalian cancer cells, to induce NF-kappaB-dependent transcripti to induce JNK/SAPK-dependent responses in mammalian cells
XX Sequence 249 AA;
Query Match 87.8%; Score 1020; DB 2; Length 249;
Best Local Similarity 88.8%; Pred. No. 2.1e-97;
Matches 199; Conservative 9; Mismatches 16; Indels 0; G
Qy 2 LSLGLALACLGLLVVSLGSWATLSAQPSQBELTAEDRRPELNPQTEESQDV
Db 26 LGLGLALACLGLLVVSLGSRASLSAQEPQAQELVAEDQPSLNPQTEESQDF
Qy 62 EQLVPRSPAPKGRKPRRAIAAHYVHPRPQDGAQAGVDGTVSGWEEKINS
Db 86 NRLVPRSPAPKGRKTRARRAIAAHYVHPRPQDGAQAGVDGTVSGWEEKINS
Qy 122 YDROIQIGFTVIRAGLYLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLEEFSA
Db 146 YNRQIGFEFTVIRAGLYLYLYCQVHFDEGKAVYKLDLLVNGVLALRCLEEFSA
Qy 182 QLRICQVSGLLPLRPGSSLRIRITLPWAHLKAAPFLTYFGLFQVH 225
Db 206 QLRICQVSGLLALRPGSSLRIRITLPWAHLKAAPFLTYFGLFQVH 249
RESULT 7
AAY95338
ID AAY95338 standard; protein; 249 AA.
AC AAY95338;
XX 25-SEP-2000 (first entry)
DE Human PRO207 antitumour protein.
XX PRO207; human; antitumour; tumour; therapy; cytostatic; breast c
KW ovarian cancer; renal cancer; colorectal cancer; uterine cancer;
KW prostate cancer; lung cancer; bladder cancer;
KW central nervous system cancer; melanoma; leukaemia; neoplasm.
OS Homo sapiens.
XX Key Location/Qualifiers
FH Peptide 1..40
FT /label= Signal_peptide
FT Modified-site 10..14
FT /note= "amidation"
FT Peptide 24..35
FT /note= "prokaryotic membrane lipoprotein lipid"
FT Modified-site 27..33
FT /note= "N-myristoylation"
FT Modified-site 29..35
FT /note= "N-myristoylation"
FT Modified-site 36..42
FT /note= "N-myristoylation"
FT Protein 41..249

IGEFIVIRAGLYLYCQVHFDEGKAVYLKDLLVDGVLAURCLBEFSATAASSIGP 205
 CQVSGLLPLRPGSSLRIRTLTPWAHLKAAPFLTYFGLFQVH 225
 CQVSGLLALRPGSSLRIRTLTPWAHLKAAPFLTYFGLFQVH 249

dard; protein; 249 AA.

(first entry)

TNF related endothelium proliferative agent).

necrosis factor; TNF; angiogenesis; wound healing; TREPA;
 ndothelium proliferative agent; tumour; metastasis;
 nerary.

Location/Qualifiers

98..249

/label= Extracellular_domain

98US-00105343.

97US-00798692.

98US-00021706.

T LAB.

760/29.

350.

ogenesis in mammal at desired sites for promoting wound
 administering soluble fragment of extracellular domain of
 s factor related endothelium proliferative agent protein.

75-76; 53pp; English.

vention relates to extracellular signal molecules,
 members of tumour necrosis factor (TNF) family molecules
 TREPA (TNF related endothelium proliferative agent).
 gically active TREPA are used to treat TREPA-associated
 ours or metastases. TREPA is used for inducing angiogenesis
 promoting wound healing and for vascularising grafted tissue
 l grafting and to promote tissue grafts. The present amino
 is clone ID #690050 human TREPA

AA;

arity 87.8%; Score 1020; DB 4; Length 249;

conservative 9; Mismatches 16; Indels 0; Gaps 0;

HALACGLLVVSLGSWATLSAQEPSOBLTAEDRPPPELNPOTESSQDWPFL 61

HALACGLLVVSLGSRASLSAQEPAPAEELVAEDQPSLNPOTESSQDPAFL 85

TPRRSAPKGRKARRPRAIAAHVHPRPQDGAQGVDTGTVSGWEETKINSSPLR 121

TPRRSAPKGRKTRARRIAAHVHPRPQDGAQGVDTGTVSGWEERINSSPLR 145

IGEFIVIRAGLYLYCQVHFDEGKAVYLKDLLVDGVLAURCLBEFSATAASSPGP 181

Db 146 YNRQIGFIVIRAGLYLYCQVHFDEGKAVYLKDLLVDGVLAURCLBEFSATAAS
 Qy 182 QLRCCQVSGLLPLRPGSSLRIRTLTPWAHLKAAPFLTYFGLFQVH 225
 Db 206 QLRCCQVSGLLALRPGSSLRIRTLTPWAHLKAAPFLTYFGLFQVH 249

RESULT 10

AAU86129

ID AAU86129 standard; protein; 249 AA.

XX AC AAU86129;

DT 15-JUL-2002 (first entry)

DE Human PRO207 polypeptide.

XX Human; PRO; benign tumour; malignant tumour; lymphoid malignancy;
 KW leukaemia; neuronal disorder; stromal disorder; blastocoeleic disc
 KW inflammatory disorder; immune disorder; angiogenic disorder; cyt
 KW neuroprotective.

XX OS Homo sapiens.

XX FN WO200153486-A1.

XX PD 26-JUL-2001.

XX PF 11-FEB-2000; 2000WO-US003565.

XX PR 08-MAR-1999; 99WO-US005028.

XX PR 11-MAR-1999; 99US-0123972P.

XX PR 11-MAY-1999; 99US-0133459P.

XX PR 02-JUN-1999; 99WO-US012252.

XX PR 22-JUN-1999; 99US-0140650P.

XX PR 22-JUN-1999; 99US-0140653P.

XX PR 20-JUL-1999; 99US-0144758P.

XX PR 26-JUL-1999; 99US-0145698P.

XX PR 17-AUG-1999; 99US-0146222P.

XX PR 31-AUG-1999; 99US-0151689P.

XX PR 01-SEP-1999; 99WO-US020111.

XX PR 15-SEP-1999; 99WO-US021090.

XX PR 30-NOV-1999; 99WO-US028313.

XX PR 01-DEC-1999; 99WO-US028301.

XX PR 01-DEC-1999; 99WO-US028634.

XX PR 05-JAN-2000; 2000WO-US000219.

XX (GETH) GENENTECH INC.

PA Ashkenazi AJ, Goddard A, Godowski PJ, Gurney AL, Hillan KJ;

PI Marsters SA, Pan J, Pitti RM, Roy MA, Smith V, Stone DM;

PI Watanabe CK, Wood WI;

XX WPI; 2002-205567/26.

XX N-PSDB; ABK40255.

XX Thirty five nucleic acids encoding PRO polypeptides, useful for t

XX benign or malignant tumors, leukemias and lymphoid malignancies,

XX inflammatory, angiogenic and immunologic disorders.

XX Claim 61; Fig 4; 302pp; English.

XX The present invention relates to the isolation of novel human PRO

XX polypeptides and the polynucleotide sequences encoding them. The

XX polypeptides, agonists, antagonists or anti-PRO antibodies are us

XX treating benign or malignant tumours (e.g. renal, kidney, bladder;

XX breast, etc), leukaemias and lymphoid malignancies, other disorde

XX as neuronal, glial, astrocytal, hypothalamic, glandular, macroph

XX stromal and blastocoeleic disorders, inflammatory, immune and angi

XX disorders. The polynucleotide sequences are also useful in gene t

XX AAU86128-AAU86162 represent the human PRO polypeptides of the in

A; 87.8%; Score 1020; DB 5; Length 249;
 rity 88.8%; Pred. No. 2.1e-97;
 nservative 9; Mismatches 16; Indels 0; Gaps 0;
 ALACIGLLLVVSLGSWATLSAQEPSQBELTAEDRRPEELNPTQESQDVPEL 61
 ALACIGLLLVVSLGSRASLSAQEPAPQELVAEEDQDPSELNPTQESQDPPEL 85
 PRRAPKGRKARPRRAIAAHYEVHPRPGDGAQAGVDTVSGWEETKINSSPLR 121
 PRRAPKGRKTRARRAIAAHYEVHPRPGDGAQAGVDTVSGWEETKINSSPLR 145
 GEFTVIRAGLYLYCQVHFDGKAVYLLKDLLVNGVLALRCLEBFSATAASS 181
 GEFIVTRAGLYLYCQVHFDGKAVYLLKDLLVNGVLALRCLEBFSATAASS 205
 QVSGLLPLRPGSSLRIRTLPAWHLKAAPFLTYFGLFQVH 225
 QVSGLLALRPGSSLRIRTLPAWHLKAAPFLTYFGLFQVH 249

(ard; protein; 249 AA.

first entry)

otein.

tumour necrosis factor; ligand; cytostatic;
 r; osteopathic.

2.

002WO-US023782.

001US-0307838P.

GENOME SCI INC.

osen CA;

59/40.

01.

imeric complex having a first polypeptide member of the
 factor (TNF) ligand family, and a second different member
 family, useful for treating cancer, osteoporosis or an
 ease.

ge 368-369; 388pp; English.

quence is the protein sequence for human TWEAK protein. The
 tes to compositions comprising heterotrimeric complexes of
 s factor (TNF) ligand family members, and their use in the
 vention and treatment of disease. In one embodiment, the
 : complex comprises full-length or extracellular portions of
 -length or extracellular portions of other TNF ligand
 i, preferably VEGI or VEGI-SV. The heterotrimeric complexes
 on are useful for treating an autoimmune disease, cancer or
 and particularly for inhibiting cancer cell proliferation,
 all proliferation, or inducing apoptosis of T cells

A;

Query Match 87.8%; Score 1020; DB 6; Length 249;
 Best Local Similarity 88.8%; Pred. No. 2.1e-97;
 Matches 199; Conservative 9; Mismatches 16; Indels 0; Gaps 0;
 QY 2 LSLGIALACIGLLLVVSLGSWATLSAQEPSQBELTAEDRRPEELNPTQESQDV 61
 Db 26 LGIGLALACIGLLLVVSLGSRASLSAQEPAPQELVAEEDQDPSELNPTQESQDP 85
 QY 62 EOLVPRRSAPKGRKARPRRAIAAHYEVHPRPGDGAQAGVDTVSGWEETKINSS 121
 Db 86 NKLIVPRRSAPKGRKTRARRAIAAHYEVHPRPGDGAQAGVDTVSGWEETKINSS 145
 QY 122 YDRQIGEFIVTRAGLYLYCQVHFDGKAVYLLKDLLVNGVLALRCLEBFSATAASS 181
 Db 146 YNRQIGEFIVTRAGLYLYCQVHFDGKAVYLLKDLLVNGVLALRCLEBFSATAASS 205
 QY 182 QLRQVSGLLPLRPGSSLRIRTLPAWHLKAAPFLTYFGLFQVH 225
 Db 206 QLRQVSGLLALRPGSSLRIRTLPAWHLKAAPFLTYFGLFQVH 249

RESULT 12

ADC35206

ID ADC35206 standard; protein; 249 AA.

XX ADC35206;

AC ADC35206;

DT 18-DEC-2003 (first entry)

DE Human TNF ligand family member #12.

XX human; tumour necrosis factor; TNF ligand; endokine alpha;
 KW excessive bone resorption disorder; osteoporosis; Paget's disease.
 KW arterial calcification.

XX Homo sapiens.

XX US2003100074-A1.

XX 29-MAY-2003.

XX 15-AUG-2002; 2002US-00218547.

XX 16-AUG-2001; 2001US-0312542P.

PR 30-OCT-2001; 2001US-0330761P.

XX (YUGG/) YU G.

PA (NIJ/) NI J.

PA (ROSE/) ROSEN C A.

PA (NARD/) NARDELLI B.

XX Yu G, Ni J, Rosen CA, Nardelli B;

PI

XX WPI; 2003-696072/66.

DR N-PSDB; ADC35205.

XX

PT New Endokine alpha gene useful for preparing a composition for treat-

PT disease associated with excessive or insufficient bone resorption

PT osteoporosis, Paget's disease or arterial calcification.

XX Disclosure; SEQ ID NO 24; 145pp; English.

PS

XX The invention relates to an isolated nucleic acid molecule encoding

CC tumour necrosis factor family ligand. A composition comprising the

CC isolated antibody or its fragment is used for treating an individual

CC need of decreased level of endokine alpha activity. The endokine

CC polypeptide present in a heterotrimeric complex is used for treat-

CC individual having a disorder associated with excessive bone resor-

CC e.g. osteoporosis, Paget's disease or arterial calcification. Tre-

CC individual having a disorder associated with insufficient bone res-

CC comprises administering an endokine alpha antagonist, which is the

CC antibody that binds specifically to endokine alpha polypeptide. T-

ence represents the amino acid sequence of a tumour necrosis factor related ligand.

AA;

Identity 87.8%; Score 1020; DB 7; Length 249;
Conservative 9; Mismatches 16; Indels 0; Gaps 0;

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 61

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 85

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 121

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 145

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 181

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 205

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 225

ELALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDVVPEL 249

Standard; protein; 284 AA.

(first entry)

tumour necrosis factor related ligand (TRELL).

neurosis factor related ligand; tnf; treatment; cancer;
sease; immune system; stimulation; suppression;
OH.

97WO-US013945.

96US-0023541P.

96US-0028515P.

97US-0040820P.

EN INC.
GENEVA FACULTY MEDICINE.

ie Y, Browning JL;

619/13.

600.

sis factor related ligand - useful for, e.g. treating cancer,
disease and immune responses to tissue grafts.

je 50-51; 69pp; English.

is that of human tumour necrosis factor related ligand
or active fragments can be included with a carrier in
al compositions to treat cancer, autoimmune diseases or
uses to tissue grafts, or to stimulate or suppress the immune
system useful to screen for TRELL receptors, by labelling with a
label and screening compositions for binding. Agents
with TRELL-receptor binding can also be screened for, can
be administered, optionally with interferon- gamma, to induce cell

death or treat, suppress or alter immune responses (especially in
human adenocarcinoma cells) involving a signal pathway between T
its receptor. It's coding sequence can be used in gene therapy f
related disorders in mammals (especially humans), e.g. tumours,
autoimmune and inflammatory diseases or inherited genetic disord
introducing into cells, and expressing, therapeutically effectiv
of a vector, e.g. a virus comprising a gene encoding TRELL. It m
be of use in the preparation of prepare probes for screening
natural/synthetic DNAs for TRELL-encoding sequences and for anti
therapy

Sequence 284 AA;

Query Match 87.8%; Score 1020; DB 2; Length 284;

Best Local Similarity 88.8%; Pred. No. 2.6e-97;

Matches 199; Conservative 9; Mismatches 16; Indels 0;

QY 2 LSLGALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDV

Db 61 LSLGALACLGILLVVSIGSWATLSAQEPSELTAEEDRRPELNPQTESQDV

QY 62 EQLVPRPSAPKGRKARRAIAAHYEVHPRPGDGAQAGVDGTVSGWEEFKINS

Db 121 NRVLPRESAPKGRKARRAIAAHYEVHPRPGDGAQAGVDGTVSGWEEFKINS

QY 122 YDQIGFTVIRAGLYLYLCVHFDEGKAVYKLDLLVNGVLALRCLEFSATAA

Db 181 YNRQIGFTVIRAGLYLYLCVHFDEGKAVYKLDLLVNGVLALRCLEFSATAA

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Db 241 QLRLCQVSGLLALRPGSSLRIRTLPAHLKAAPFLTYFGLFOVH 284

RESULT 14

AAW93590

ID AAW93590 standard; protein; 208 AA.

XX AC AAW93590;

XX DT 18-JUN-1999 (first entry)

XX DE Human TNRL3 protein.

XX KW Tumour necrosis factor receptor; signal transducer molecule; TNF

XX KW developmental abnormality; Gestational abnormality; prostate c

XX KW APO6; APO8; APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy;

XX KW cytoplasmic domain; immunogen; antibody preparation; breast carc

XX OS Homo sapiens.

XX PN WO9911791-A2.

XX PD 11-MAR-1999.

XX PF 04-SEP-1998; 98WO-US018393.

XX PR 05-SEP-1997; 97US-00924634.

XX PA (UNIW) UNIV WASHINGTON.

XX PI Chaudhary PM;

XX DR WPI; 1999-205191/17.

XX DR N-PSDB; AAX23424.

XX PT New Tumor Necrosis Factor family receptor polypeptides and ligand

XX PT useful for diagnosis and treatment of prostate cancer and develop

XX PS Claim 40; Fig 13A; 156pp; English.

GenCore version 5.1.6
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ic search, using sw model

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(without alignments)

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-09-245-198A-3

73

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aximum Match 100%

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the number of results predicted by chance to have a
than or equal to the score of the result being printed,
ad by analysis of the total score distribution.

SUMMARIES

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1.9	948	13	BQ707185	BQ707185 AGENCOURT
1.4	963	13	BQ671259	BQ671259 AGENCOURT

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6	725.4	52.8	1071	12	BM921213	€
7	701.4	51.1	731	12	BI871711	€
8	688	50.1	828	12	BI596681	€
9	677.4	49.3	728	12	BI870393	€
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IMAGE:154742, mRNA sequence.
ACCESSION
BX090012
VERSION
BX090012.1 GI:27821952
KEYWORDS
EST.
SOURCE
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ORGANISM
Homo sapiens
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 776)
AUTHORS
Bert, L., Heil, O., Hennig, S., Neubert, P., Patsch, E., Pete
Radelof, U., Schneider, D. and Korn, B.
TITLE
Human UnigeneSet - RZPD3
JOURNAL
Unpublished (2003)
COMMENT
Contact: Ina Rofls
RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany
RZPD; IMAGP998E15243.
RZPDLIB; I.M.A.G.E. cDNA Clone Collection;
Human UnigeneSet - RZPD3 (RZPDLIB No.972)
http://www.rzpd.de/CloneCards/cgi-
bin/showlib.pl/cgi/response?libNo=972 Contact: Ina Rofls
RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
Heubnerweg 6, D-14059 Berlin, Germany

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DEFINITION	BI766766	mRNA sequence.	
ACCESSION	BI766766	BI766766.1 GI:15758344	
VERSION	EST.		
KEYWORDS	Homo sapiens (human)		
SOURCE	Homo sapiens		
ORGANISM			
REFERENCE			
AUTHORS	1	(Bases 1 to 834)	
TITLE	National Institutes of Health, Mammalian Gene Collection		
JOURNAL	Unpublished (1999)		
COMMENT	Contact: Robert Strausberg, Ph.D. Email: cgapbs-remail.nih.gov Tissue Procurement: Life Technologies, Inc. cdna Library Preparation: Life Technologies, Inc. cdna Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Incyte Genomics, Inc. Clone distribution: MGC clone distribution information c found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: LLAM1517 row: c column: 18 High quality sequence stop: 772.		
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Matches	818; Conservative	0; Mismatches 12; Indels 5;	
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 5.1 GI:21846084
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 ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 es 1 to 948)
 http://mgi.nci.nih.gov/
 1 Institutes of Health, Mammalian Gene Collection (MGC)
 shed (1999)
 : Robert Strausberg, Ph.D.
 cgabs-r@mail.nih.gov
 Procurement: Dr. Mark Watson
 Library Preparation: Rubin Laboratory
 Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 quencing by: Agencourt Bioscience Corporation
 distribution: MGC clone distribution information can be
 through the I.M.A.G.E. Consortium/LLNL at:
 image.llnl.gov
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 laboratory of Gerald M. Rubin (University of Cal
 Berkeley) using ZAP-cDNA synthesis kit (Stratagen
 Superscript II RT (Life Technologies). Note: this
 NIH_MGC Library."
 ORIGIN
 Query Match 54.9%; Score 753.2; DB 13; Length 948;
 Best Local Similarity 95.8%; Pred. No. 1.8e-131;
 Matches 807; Conservative 0; Mismatches 30; Indels 7; G
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 DB 1 ACCGTCGGAACCTGAATCCCGACAGAGAGANAGCCAGGATCCTGCGCCTTCTCTG
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 QY 425 TCGACGCCCATTTATGAAGTTTCATCCAGACCTGGACAGGACCGGAGCGCAGGAGT
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ORIGIN

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apiens
apiens

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uses 1 to 731)

!C <http://mgc.nci.nih.gov/>.

National Institutes of Health, Mammalian Gene Collection (MGC)

ished (1999)

Dr. Robert Strausberg, Ph.D.

cgapbs-r@mail.nih.gov

: Procurement: ATCC

Library Preparation: Life Technologies, Inc.

Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information C:
found through the I.M.A.G.E. Consortium/LLNL at:
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69. The seventieth column is labeled "SOURCE".	
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71. The seventy-second column is labeled "SOURCE".	
72. The seventy-third column is labeled "FEATURES".	
73. The seventy-fourth column is labeled "SOURCE".	
74. The seventy-fifth column is labeled "FEATURES".	
75. The seventy-sixth column is labeled "SOURCE".	
76. The seventy-seventh column is labeled "FEATURES".	
77. The seventy-eighth column is labeled "SOURCE".	
78. The seventy-ninth column is labeled "FEATURES".	
79. The eightieth column is labeled "SOURCE".	
80. The eighty-first column is labeled "FEATURES".	
81. The eighty-second column is labeled "SOURCE".	
82. The eighty-third column is labeled "FEATURES".	
83. The eighty-fourth column is labeled "SOURCE".	
84. The eighty-fifth column is labeled "FEATURES".	
85. The eighty-sixth column is labeled "SOURCE".	
86. The eighty-seventh column is labeled "FEATURES".	
87. The eighty-eighth column is labeled "SOURCE".	
88. The eighty-ninth column is labeled "FEATURES".	
89. The ninetieth column is labeled "SOURCE".	
90. The ninety-first column is labeled "FEATURES".	
91. The ninety-second column is labeled "SOURCE".	
92. The ninety-third column is labeled "FEATURES".	
93. The ninety-fourth column is labeled "SOURCE".	
94. The ninety-fifth column is labeled "FEATURES".	
95. The ninety-sixth column is labeled "SOURCE".	
96. The ninety-seventh column is labeled "FEATURES".	
97. The ninety-eighth column is labeled "SOURCE".	
98. The ninety-ninth column is labeled "FEATURES".	
99. The hundredth column is labeled "SOURCE".	

Location/Qualifiers
1. .731

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

```
/clone="IMAGE:5405478
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```
/tissue_type="adenocarcinoma, cell line"
```

```
/lab_host="DH10B (phage-resistant)"
```

```
/clone_lib="NIH_MGC_90"
```

```
/note="Organ: liver; Vector: pCMV-SPORT6; Site_1:
```

Site 2: SalI; Cloned unidirectionally; oligo-dT 1

Average insert size 1.7 kb. Library enriched for

full-length clones and constructed by Life Techn

Note: this is a NIH_MGC Library."

ORIGIN

Query Match	51.1%;	Score 701.4;	DB 12;	Length 731;
Best Local Similarity	99.3%;	Pred. No. 9.4e-122;		
Matches 726; Conservative	0;	Mismatches 1;	Indels 4;	


```

1  TGAACCGACTAGTTCGGCTCGCAGAAAGTGACCTAAAGCGCGAAACACACGGC 413
2  |||||
3  TGAACCGACTAGTTCGGCTCGCAGAAAGTGACCTAAAGCGCGAAACACACGGC 360
4  |||||
5  AGAGCGATCCAGCCCATTAAGTTATCCACGACCTGGACGAGCGAGCGCA 473
6  |||||
7  AGAGCGATCCAGCCCATTAAGTTATCCACGACCTGGACGAGCGAGCGCA 420
8  |||||
9  TGTGTGACGGGACAGTGAGTGGCTGGGAGGAGCCAGAAATCAACAGCTCCAGCCC 533
10 TGTGTGACGGGACAGTGAGTGGCTGGGAGGAGCCAGAAATCAACAGCTCCAGCCC 480
11 |||||
12 TGTATACCGCCAGATCGGGAGTTATAGTACACCGGGCTGGCTTACTACCT 593
13 |||||
14 TGTATACCGCCAGATCGGGAGTTATAGTACACCGGGCTGGCTTACTACCT 540
15 |||||
16 TGTAGGTGACCTTGTATGAGGGAGAGGCTGTCTACTGAAGCTGGACTTGTGGT 600
17 |||||
18 TGTGTGCTGGCCCTCGCTGCTGGAGGAATTCAGCCACTGCGGCGCAGTTCCCT 713
19 |||||
20 TGTGTGCTGGCCCTCGCTGCTGGAGG-ATTCTAGCCACTGCGGCGAGTTCCCT 659
21 |||||
22 TCCAGCTCGGCTCTGCGAGGTGTCTGGGCTGT 751
23 |||||
24 CCCAGCTCGGCTCTGCGAGGTGTCTGGGCTGT 697

```

```

19 170-avw-m-22-0-UI.r1 NIH_MGC_214 Homo sapiens cDNA clone
19 10559317 5', mRNA sequence.
19

```

```

19 9.1 GI:33203878

```

```

19 piens (human)
19 ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
19 ta; Eutheria; Primates; Catarrhini; Hominidae; Homo.
19 es 1 to 666)
19 ,M.F., Lennon,G. and Soares,M.B.
19 zation and subtraction: two approaches to facilitate gene
19 Res. 6 (9), 791-806 (1996)
19

```

```

19 : Soares, MB
19 ated Laboratory for Computational Genomics
19 ity of Iowa
19 ton Road , 4156 MEBRF, Iowa City, IA 52242, USA
19 9 335 8250
19 9 335 9565
19 bento-soares@uiowa.edu
19 Procurement: Mary Hendrix
19 library preparation: Dr. M. Bento Soares, University of Iowa
19 library Arrayed by: Dr. M. Bento Soares, University of Iowa
19 quencing by: Dr. M. Bento Soares, University of Iowa
19 Distribution: Distribution information can be found at
19 genome.uiowa.edu/distribution/humanfl.html
19 mer: pYX-5.
19 Location/Qualifiers
19 1. .666
19 /organism="Homo sapiens"
19 /mol_type="mRNA"
19 /db_xref="taxon:9606"
19 /clone="IMAGE:30559317"
19 /tissue_type="Chondrosarcoma Lung Metastasis cell lines"
19 /lab_host="DH10B (T1 phage resistant)"
19 /clone_lib="NIH_MGC_214"
19 /note="Organ: Lung; Vector: pYX-Asc; Site 1: EcoR I;
19 Site 2: Not I; The library was constructed according

```

Bonaldo, Lennon and Soares, Genome Research, 6:79
1996. Denatured RNA was size fractionated on a 1%
gel. First strand cDNA synthesis was primed with
primer containing a Not I site. Double strand cDN
size selected according to mRNA size fraction, li
with EcoR I adaptor, digested with Not I and then
directionally into pYX-Asc vector. The library ta
sequence located between the Not I site and the p
is GATAAGGCCA. Tissue was provided by Mary Hendri

ORIGIN

```

Query Match 47.8%; Score 656.6; DB 14; Length 666;
Best Local Similarity 98.9%; Pred. No. 2.5e-113;
Matches 659; Conservative 0; Mismatches 7; Indels 0; G
292 GAGGAGGACGAGGACCGCTCGGAACCTGAATCCCGACAGAGAAAGCCAGGATCC
Db 1 GAGGAGGACCGGACCGCTCGGAACCTGAATCCCGACAGAGAAAGCCAGGATCC
352 CCTTTCTGAAACCGACTAGTTCGGCTCGCAGAAAGTGACCTAAAGCGCGAAAC
Db 61 CCTTTCTGAAACCGACTAGTTCGGCTCGCAGAAAGTGACCTAAAGCGCGAAAC
412 GCTCGAAGAGCGATCGAGCCCATTTATGAAGTTTCATCCACGACCTGGACAGGCGG
Db 121 GCTCGAAGAGCGATCGAGCCCATTTATGAAGTTTCATCCACGACCTGGACAGGCGG
472 CAGGCAGGTGTGACGCGGACAGTGAGTGGCTGGGAGGAGCCAGAAATCAACAGCTC
Db 181 CAGGCAGGTGTGACGCGGACAGTGAGTGGCTGGGAGGAGCCAGAAATCAACAGCTC
532 CTTCTCGGCTACAAACCGCAGATCGGGGAGTTTATAGTCACCCGGGCTGGGCTCTA
Db 241 CTTCTCGGCTACAAACCGCAGATCGGGGAGTTTATAGTCACCCGGGCTGGGCTCTA
592 CTGTACTGTCAAGTGCACCTTTGATGAGGGAAGGCTGTCTACCTGAAGCTGGACTT
Db 301 CTGTACTGTCAAGTGCACCTTTGATGAGGGAAGGCTGTCTACCTGAAGCTGGACTT
652 GTGGATGTTGCTGTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCG
Db 361 GTGGATGTTGCTGTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCGCTCG
712 CTCGGGCCCCAGCTCCGCTCTGCCAGGTGTCTGGGCTGTGTGGCTGTGTGGCTGTGT
Db 421 CTCGGGCCCCAGCTCCGCTCTGCCAGGTGTCTGGGCTGTGTGGCTGTGTGGCTGTGT
772 TCCTTCGGATCGGACACCTCCCTCGCTGGGCCCCATCTCAAGGCTGCCCCCTTCCTCAC
Db 481 TCCTTCGGATCGGACACCTCCCTCGCTGGGCCCCATCTCAAGGCTGCCCCCTTCCTCAC
832 TTTCGGACTCTTCAGGTTTCACTGAGGGGCCCCCTGGTCTCCCGACAGTGTGTCAGGCT
Db 541 TTTCGGACTCTTCAGGTTTCACTGAGGGGCCCCCTGGTCTCCCGACAGTGTGTCAGGCT
892 GGCTCCCCCTCGACAGCTCTCTGGGACACCGGTCCTCTGCCCCCAGCCCTCAGCCGCT
Db 601 GGCTCCCCCTCGACAGCTCTCTGGGACACCGGTCCTCTGCCCCCAGCCCTCAGCCGCT
952 TGCTCC 957
Db 661 TGCTCC 666

```

```

RESULT 12
Bi966060/c
LOCUS
DEFINITION
ie72g04.x1 Melton Normalized Human Islet 4 N4-HIS 1 Homo sa
cDNA clone IMAGE:5672623 3' similar to FR:054907 054907 TNF
WEAK INDUCER OF APOPTOSIS ;, mRNA sequence.
ACCESSION
VERSION
KEYWORDS
EST.

```

QY	1027	TTCCCACTCTTATCTTAAACTCCCCCACCGCCCACTCTCCACCTCAGTAGTCCTCC
Db	369	TTCCCACTCTTATCTTAAACTCCCCCACCGCCCACTCTCCACCTCAGTAGTCCTCC
QY	1087	CCGTGACCCTTTGAGGCCCCAGTGATCTCGATCTCCCCCCTGGGCACAGACCCCC
Db	309	CCCTGACCCTTTGAGGCCCCAGTGATCTCGATCTCCCCCCTGGGCCACAGACCCCC
QY	1147	ATTGTGTTCACTGTACTCTGTGGGCAAGGATGGTCCAGAAGACCCCACTTCAGG
Db	249	ATTGTGTTCACTGTACTCTGTGGGCAAGGATGGTCCAGAAGACCCCACTTCAGG
QY	1207	AGAGGGGCTGACCTGGCGGCGAGGAAGCAAAGAGACTGGGGCTTAGGCAGGAGT
Db	189	AGAGGGGCTGACCTGGCGGCGAGGAAGCAAAGAGACTGGGGCTTAGGCAGGAGT
QY	1267	AATGTGAGGGCGGAGAAACAAGACAGCTCCTCCCTTGAGAAATTCCTGTGGATT
Db	129	AATGTGAGGGCGGAGAAACAAGACAGCTCCTCCCTTGAGAAATTCCTGTGGATT
QY	1327	AACGATATTAATTTTTATTATTATTTGACAAAATGTTGATAAATGG 1373
Db	69	AACGATATTAATTTTTATTATTATTTGACAAAATGTTGATAAATGG 23
RESULT 13		
AF163779 Human Homo sapiens genomic clone BAC750E14, genor		
LOCUS	AF163779	1027 bp DNA linear GSS 2:
DEFINITION	sequence.	
ACCESSION	AF163779	
VERSION	GI:5726439	
KEYWORDS	GSS.	
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele;	
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo	
TITLE	Cousin,P., Billette,J., Chaubert,P. and Shaw,P.H.	
JOURNAL	Physical map of 17p13 and the genes adjacent to p53	
MEDLINE	Genomics 63 (1), 60-68 (2000)	
PUBMED	20130114	
COMMENT	Contact : Shaw PH Experimental Oncology Institute of Pathology Rue du Bugnon 25, Lausanne, VD 1011, Switzerland sub clone=AB2R Asc-BamHI PSL1180 Class: BAC subclone. Location/Qualifiers 1..1027	
FEATURES	/organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" /map="17p" /clone="BAC750E14" /clone_lib="Human"	
ORIGIN		
Query Match 42.3%; Score 581.4; DB 28; Length 1027;		
Best Local Similarity 98.0%; Pred. No. 4.le-99;		
Matches 577; Conservative 8; Mismatches 4; Indels 0;		
QY	785	GCACCTTCCCTGGGCCCATCTCAAGGCTGCCCCCTTCTCTCACTTCTCGACT
Db	1	GCWCCCTCCCTGGGCCCATCTCAAGGCTDYVCCCTTCTCTCACTTCTCGACT
QY	845	AGGTTCACTGAGGGCCCTGGTCTCCCAACAGTCGTCCAGGCTGCCGCTCCCTCC
Db	61	AGGTTCACTGAGGGCCCTGGTCTCCCAACAGTCGTCCAGGCTGCCGCTCCCTCC
QY	905	AGCTCTCTGGGSCACCGGTCCTCTGCCCCACCTCTCAGCGCTCTTTTGTCTCAG

```

|||||
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|||||
CCCTCTAGAGCTGCTGGGCTGTTCACGTTTTCATCCACATAAATACAG 1024
|||||
CCCTCTAGAGCTGCTGGGCTGTTCACGTTTTCATCCACATAAATACAG 240
|||||
CCACTCTATCTTAACTCCCAACCCGACCTCTCCACCTCACTAGTCCCA 1084
|||||
CCACTCTATCTTAACTCCCAACCCGACCTCTCCACCTCACTAGTCCCA 300
|||||
TGACCTTTGAGGCCCCAGTGTCTGACTCCCTCCCTGGCCACAGACCCCA 1144
|||||
TGACCTTTGAGGCCCCAGTGTCTGACTCCCTCCCTGGCCACAGACCCCA 360
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|||||
TGCTTCTACTGTACTGTGGGCAAGGATGGTCCAGAGACCCCACTTCCAG 420
|||||
GGGCTGACCTTGGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGTCC 1264
|||||
GGGCTGACCTTGGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGTCC 480
|||||
GTGAGGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGTCC 1324
|||||
GTGAGGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGTCC 540
|||||
AGATATATTTTATTTATTTATTTATTTATTTATTTATTTATTTATTT 1373
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AGATATATTTTATTTATTTATTTATTTATTTATTTATTTATTTATTT 589

```

```

9 567 bp mRNA linear EST 06-NOV-2002
.y1 Human Retinal pigment epithelium/choroid cDNA
malized, unamplified): cs Homo sapiens cDNA clone cs80h07
A sequence.
9.1 GI:24733297
piens (human)
piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 567)
n,J.W., Bouffard,G., Smith,D. and Peterson,K.
ed sequence tag analysis of human RPE/choroid for the
Project: Over 6000 non-redundant transcripts, novel genes
ice variants
s. 8 (4), 205-220 (2002)
0
0
: Wistow G
on Molecular Structure and Function
l Eye Institute
NIH, Bethesda, MD 20892-2740, USA
1 402 3452
1 496 0078
graeme@helix.nih.gov
80 row: h column: 07
mer: M13RPI reverse primer (ABI).
Location/Qualifiers
1. .567
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="cs80h07"
/tissue_type="RPE/choroid"
/dev_stage="Adult"
/lab_host="EMDH10B"
/clone_lib="Human Retinal pigment epithelium/choroid cDNA

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RESULT 15
BO884231
LOCUS
DEFINITION
IMAGE:6197488 5', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutelec
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

```

(Un-normalized, unamplified): cs
/notes="Organ: Eye; Vector: pCMVSPORT6; Two differ
eyes (75-80 years old) yielded approximately 600
dissected RPE/choroid tissue. This in turn yield
of total RNA and 7 ug of mRNA. A directionally cl
library in the pCMVSPORT6 vector was constructed
Technologies (Rockville, MD; now part of Invitro
essentially following the protocols of the SuperS
Plasmid System (Invitrogen Corp.
http://www.invitrogen.com/>). The library code
designation was cs. For this library, cDNA insert
cloned into the NotI/MluI sites of the vector. ES
analysis was performed on the unamplified library
NIH Intramural Sequencing Center (MISC)."

ORIGIN

Query Match	Best Local Similarity	Score	DB 14;	Length	567;
Matches	567;	Conservative	0;	Mismatches	0;
Indels	0;	Gt			
55	CTCGGCTCCCGGATGGGGGGGGGTGAGGACGACACGCCCCCGCCCATGGG				
1	CTCGGCTCCCGGATGGGGGGGGGTGAGGACGACACGCCCCCGCCCATGGG				
115	CGTCGGAGCCAGAGCGAGGGGGGGCGCCGGGGGGAGCCGGGCGACCCGCTGTGGT				
61	CGTCGGAGCCAGAGCGAGGGGGGGCGCCGGGGGGAGCCGGGCGACCCGCTGTGGT				
175	CTCGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGG				
121	CTCGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGG				
235	TTGGGGAGCCGGGCGATCGCTGTCGCCCGGAGGAGCTGCCCGAGGAGGCTGTGGG				
181	TTGGGGAGCCGGGCGATCGCTGTCGCCCGGAGGAGCTGCCCGAGGAGGCTGTGGG				
295	GAGGACCCAGAGCCCGCTCGGAATGAATCCCGAGAGAGAAAGCCAGGATCTCGC				
241	GAGGACCCAGAGCCCGCTCGGAATGAATCCCGAGAGAGAAAGCCAGGATCTCGC				
355	TTCTCTGAACCGACTAGTTTCGGCTCGAGAGTGCACCTAAAGGCCGGAACACCG				
301	TTCTCTGAACCGACTAGTTTCGGCTCGAGAGTGCACCTAAAGGCCGGAACACCG				
415	CGAAGAGCGATCGAGCCCATTTATGAAGTTTCATCCAGACCTCGGACGACCGAGG				
361	CGAAGAGCGATCGAGCCCATTTATGAAGTTTCATCCAGACCTCGGACGACCGAGG				
475	GCAGGTGTGGAGCGGACAGTGTAGTGGCTGGGAGGAGCCAGAGTCAACAGCTCCAG				
421	GCAGGTGTGGAGCGGACAGTGTAGTGGCTGGGAGGAGCCAGAGTCAACAGCTCCAG				
535	CTGGCGTCAACCGCCAGATCGGGAGTTTATAGTCAACCGGGCTGGGCTCTACTA				
481	CTGGCGTCAACCGCCAGATCGGGAGTTTATAGTCAACCGGGCTGGGCTCTACTA				
595	TACTGTGAGTGCACCTTTGATGAGGG 621				
541	TACTGTGAGTGCACCTTTGATGAGGG 567				

```

BO884231
AGENCY: 8682031
IMAGE: 6197488 5', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutelec
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

```

ases 1 to 940)
 3C http://mgc.nci.nih.gov/
 nal Institutes of Health, Mammalian Gene Collection (MGC)
 lished (1999)
 ct: Robert Strausberg, Ph.D.
 : cgsbs-f@mail.nih.gov
 e procurement: Dr. James R. Lupski
 Library Preparation: Life Technologies, Inc.
 Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 Sequencing by: Agencourt Bioscience Corporation
 e distribution: MGC clone distribution information can be
 through the I.M.A.G.E. Consortium/LNL at:
 //image.lnl.gov

: L1AM13607 row: j column: 17
 quality sequence stop: 453.
 Location/Qualifiers
 1. .940

/organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:6197488"
 /sex="male"
 /tissue_type="sciatic nerve"
 /dev_stage="adult, 70 yr"
 /lab_host="DH10B"
 /clone_lib="Lupski sciatic nerve"
 /notes="Vector: pCMV-SPORT6 (Life Technologies); Site 1:
 NotI; Site 2: SalI; cDNA made by oligo-dT priming.
 Directionally cloned using the following adaptors:
 5'-TCGACCCACGCGTCCG-3' and
 5'-GACTAGTTCAGATCGGCGCGCCCT(15)-3'. Size selected >
 1 kb for average insert length 1.87 kb. This is a primary
 library, non-amplified. Library constructed by Life
 Technologies and donated by J. Lupski, M.D./Ph.D. (Baylor
 College of Medicine) and is available through Life
 Technologies."

40.9%; Score 562; DB 13; Length 940;
 larity 96.5%; Pred. No. 1.8e-95;

Conservative 0; Mismatches 15; Indels 9; Gaps 8;

ACCGCTCGGAACCTGAATCCCGACAGAGAAAGCCAGGATCCTGGCCCTTTCCCTG 360
 ACCCGTCGGAACCTGAATCCCGACAGAGAAAGCCAGGATCCTGGCCCTTTCCCTG 60

AGACTAGTTCGGGCTCGCAGAGTGCACCTAAAGCGCGGAACACGGGCTCGAAGA 420

AGACTAGTTCGGGCTCGCAGAGTGCACCTAAAGCGCGGAACACGGGCTCGAAGA 120

ATCGAGCCCATTTATGAAGTTTCATCCAGACCTGGACAGGAGCGAGCGAGGAGGT 480

ATCGAGCCCATTTATGAAGTTTCATCCAGACCTGGACAGGAGCGAGCGAGGAGGT 180

ACGGGACAGTGAAGTGGTGGGAGGAGCCAGAAATCAACAGCTCCAGCCCTTCGGCC 540

ACGGGACAGTGAAGTGGTGGGAGGAGCCAGAAATCAACAGCTCCAGCCCTTCGGCC 240

ACCGCCAGATCGGGGAGTTTATAGTCACCGGGCTGGGCTCTACTACTGTACTGT 600

ACCGCCAGATCGGGGAGTTTATAGTCACCGGGCTGGGCTCTACTACTGTACTGT 300

ATGCACTTTGATGAGGGGAGGCTGTCTACCTGAAGCTGGACTTGTGTGGTGGATGGT 660

ATGCACTTTGATGAGGGGAGGCTGTCTACCTGAAGCTGGACTTGTGTGGTGGATGGT 360

ATGGCCCTTGGCTGCTCGGAGGAATTCCTAGCCACTGGCGGCCAGTTTCCCTTCGGGCC 720

ATGGCCCTTGGCTGCTCGGAGGAATTCCTAGCCACTGGCGGCCAGTTTCCCTTCGGGCC 420

ATCCGCTCTG-CGAGGTGTCTGGGCTGTGGCCCTGGCGCCA-GGGTCTCCCTGC 778

ATCCGCTCTGCCCCAGGTGTCTGGGCTGTGGCCCTGGCGGCCAGGGGCCCTCCCTGC 480

QY 779 GGATCCGCACCCCTCCCTCGGGCCCATCTCAAGGCTG-CCGCCCTTCTCCTACCTAC-
 Db 481 GGATCCGCACCCCTCCCTCGGGCCCATCTCAAGGCTGCCCCCTTTCTCCTACTACT
 QY 837 ACTCTTCCAGGTTTCACTGAGGGGCCCTTGGTCTCTCCCAACAGTCTGT-CCCAGGCTGC
 Db 541 ACTCTTCCAGGTTTCACTGAGGGGCCCTTGGTCTCTCCCAACAGTCTGTCTCCCAAGGCTGC
 QY 895 TCCCTCTGACAGTCTCT-GGGACACCGGTTCCCTCTGCCCCACCC--TCAGCCG
 Db 601 TCCCTCTGACAGTCTCTGAGGAAACCGGTTCCCTCTGCCCCACCCCTCAGGCGG
 QY 952 TGCTCCAGACCTGCGCCCTCCCTCT 975
 Db 661 TGGTCCAGAACTGGCCCTCCCTCT 684

Search completed: April 7, 2004, 23:15:25
 Job time : 3339.48 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

ic search, using sw model

ril 7, 2004, 17:30:20 ; Search time 5268.84 Seconds
(without alignment)
11294.703 Million cell updates/sec

-09-245-198a-3

73

atgcattgttagactttga.....gacaaaatgtgataaatgg 1373

ENTITY NUC

pop 10.0 , Gapext 1.0

70272 seqs, 21671516995 residues

ts satisfying chosen parameters: 6940544

3th: 0

3th: 2000000000

inimum Match 0%

aximum Match 100%

isting first 45 summaries

enEmbl.*

gb_ba.*

gb_htg.*

gb_in.*

gb_on.*

gb_ov.*

gb_pat.*

gb_ph.*

gb_pl.*

gb_pr.*

gb_ro.*

gb_sts.*

gb_sy.*

gb_un.*

gb_vi.*

em_ba.*

em_fun.*

em_hum.*

em_in.*

em_mu.*

em_or.*

em_ov.*

em_pat.*

em_ph.*

em_pl.*

em_ro.*

em_sts.*

em_un.*

em_vi.*

em_htg_hum.*

em_htg_inv.*

em_htg_other.*

em_htg_mus.*

em_htg_pln.*

em_htg_rnd.*

em_htg_mam.*

em_htg_vrt.*

em_sy.*

em_htgo_hum.*

em_htgo_mus.*

em_htgo_other.*

the number of results predicted by chance to have a

score greater than or equal to the score of the result being pri
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	1373	100.0	1373	6	BD062758	BD062758 A
2	1325	96.5	1421	6	BD090952	BD090952 Af
3	1320.2	96.2	1353	6	AX201324	AX201324 Se
4	1320.2	96.2	1353	9	AY358870	AY358870 Hc
5	1320.2	96.2	1368	9	AF055872	AF055872 Hc
6	1285	93.6	1306	9	AF030099	AF030099 Hc
7	1226.4	89.3	1236	6	AR140407	AR140407 Se
8	1226.4	89.3	1236	6	BD057124	BD057124 Me
9	1096.8	79.9	1642	9	BC019047	BC019047 Hc
10	768.2	56.0	60268	9	AC016876	AC016876 Hc
11	754	54.9	218485	2	AC127470	AC127470 Pa
12	683.4	49.8	1239	10	AF030100	AF030100 M
13	629.2	45.8	898	6	AX180714	AX180714 Se
14	614.6	44.8	1168	6	BD062757	BD062757 A
15	553	40.3	1816	9	AY081051	AY081051 Hc
16	364.2	26.5	148555	2	AC126921	AC126921 Bc
17	309.6	22.5	203083	2	AC069459	AC069459 Mu
18	309.6	22.5	234182	10	AL603707	AL603707 M
19	302.2	22.0	130254	2	AC136195	AC136195 Ra
20	302.2	22.0	165316	2	AC119115	AC119115 Ra
21	302.2	22.0	223877	2	AC098923	AC098923 Ra
22	302.2	22.0	225077	2	AC136563	AC136563 Ra
23	283.4	20.6	149736	2	AC126239	AC126239 Fe
24	267	19.4	234801	2	AC118309	AC118309 Ra
25	246.4	17.9	180222	2	AC130192	AC130192 Su
26	242.4	17.7	176258	2	AC126925	AC126925 Ca
27	193.4	14.1	195	6	AX379024	AX379024 Se
28	184.2	13.4	212093	2	AC126237	AC126237 Ca
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RESULT 1	BD062758	1373 bp	DNA	linear	PAT 27
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DEFINITION	A tumor necrosis factor related ligand.				
ACCESSION	BD062758				
VERSION	BD062758.1	GI:22608361			
KEYWORDS	JP 2001505407-A/2.				
SOURCE	unidentified				
ORGANISM	unclassified				
REFERENCE	1 (bases 1 to 1373)				
AUTHORS	Chicheportiche, Y. and Browning, J. L.				
TITLE	A tumor necrosis factor related ligand				
JOURNAL	Patent: JP 2001505407-A 2 24-APR-2001;				
	BIOTEN INC, THE FACULTY OF MEDICINE OF THE UNIVERSITY OF GE				

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ACCESSION		BD090952				
VERSION		BD090952.1	GI:22636562			
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ORGANISM		Homo sapiens				
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AUTHORS		1 (Bases 1 to 1421)				
TITLE		Askenazi, A. J., Marsters, S. A. and Pitti, R.				
JOURNAL		Apo-3 ligand polypeptide				
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Location/Qualifiers

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 Goddard,A., Wood,W.I. and Godowski,P.
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 MGC help desk
 gapbs-x@mail.nih.gov
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 burg, Maryland;

Web site: <http://www.nisc.nih.gov/>
 Contact: nisc_mgcnhgri.nih.gov
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DEFINITION	Homo sapiens chromosome 17, clone RP11-186B7, complete se
ACCESSION	AC016876
VERSION	AC016876.10
KEYWORDS	HTG, GI:24431829
SOURCE	Homo sapiens (human)
ORGANISM	Homo sapiens

REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS

1 (bases 1 to 60268)
Birren, B., Nusbaum, C. and Lander, E.
Mammalia; eutheria; Primates; Catarrhini; Hominidae; Homo
Unpublished

2 (bases 1 to 60268)
Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Ainc
Baldwin, J., Barna, N., Becker, R., Boguslavsky, L., Bouckge
Brown, A., Castle, A., Colangelo, M., Collins, S., Collymore, J.
Cooke, P., Dearellano, K., Dewar, K., Domingo, M., Donelan, L.,
Ferreira, P., Fitzhugh, W., Forrest, C., Funke, R., Gage, D.,
Galagan, J., Gardyna, S., Grant, G., Hagos, B., Heaford, A., Hc
Howland, J. C., Johnson, R., Jones, C., Kann, L., Karatas, A., K
Lehoczky, J., Lieu, C., Locke, K., Macdonald, P., Marquis, N.,
McSwan, P., McGurk, A., McKernan, K., McLaughlin, J., Melidrm,
Morrow, J., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell,
Peterson, K., Pollara, V., Riley, R., Roy, A., Santos, R., Seve
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas,
Tsfaye, S., Tirrell, A., Vassiliev, H., Vo, A., Wheeler, J., W
Wyman, D., Ye, Y., Zimmer, A. and Zody, M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Wyman, D., de, W. J., Zimmer, A. and Zody, M.
Direct Submission
Submitted (08-DEC-1999) Whitehead Institute/MIT Center for
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 60268)
Birren, B., Nuebaum, C., Lander, E., Ali, A., Allen, N., Anders
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalgalte
Camarata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A.
Cook, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J. S., Dodi
Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat
Karatas, A., Kells, C., Lander, T., Levine, R., Lindblad-Toh,
Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C.,
McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mienga, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norma,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, I.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Ro
Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Sevi
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, M., Ta
Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, I.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zaint
Zembek, L., Zimmer, A. and Zody, M.

JOURNAL
REFERENCE
AUTHORS

Submitted (08-OCT-2002) Whitehead Institute/MIT Center for
Research, 320 Charles Street, Cambridge, MA 02141, USA
4. (bases 1 to 60268)
Birren,B., Nusbaum,C., Lander,E., Ali,A., Allien,N., Andersc
Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
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Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galaagan,J.,
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A., Karatas, A., Kells, C., Landers, T., Levine, R.,
 ad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J.,
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 V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,
 C.H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J.,
 on, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R.,
 Rogov, P., Roman, J., Roy, A., Schauer, S., Schupbach, R.,
 S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,
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 S.M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X.,
 D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
 Submission
 ted (31-OCT-2002) Whitehead Institute/MIT Center for Genome
 ch, 320 Charles Street, Cambridge, MA 02141, USA
 31, 2002 this sequence version replaced gi:233592141.
 peats were identified using RepeatMasker:
 A.F.A. & Green, P. (1996-1997)
 /ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 nter: Whitehead Institute/ MIT Center for Genome Research
 nter code: WIBR
 o site: http://www-seq.wi.mit.edu
 ntract: sequence submissions@genome.wi.mit.edu
 ----- Project Information
 iter project name: L3849
 iter clone name: 186_B_7

ie first 60.3 kilobases of this clone are being submitted.
 nander overlaps accession number AC113189 [WICGR project

Location/Qualifiers

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 24279..24530
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 Matches 770; Conservative 0; Mismatches 3; Indels 0; Gs

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Abstract The purpose of this study was to determine whether there were differences in the prevalence of risk factors for coronary artery disease between men who had been exposed to asbestos and those who had not. A case-control study was conducted among men aged 60 years or older who had been employed in asbestos-related occupations for at least 10 years. The prevalence of risk factors for coronary artery disease was compared between cases (men with a history of myocardial infarction) and controls (men without a history of myocardial infarction). The results showed that the prevalence of risk factors for coronary artery disease was significantly higher in cases than in controls. This suggests that exposure to asbestos may be associated with an increased risk of developing coronary artery disease.

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25 499 bp DNA PAT 18-DEC-2003
 25 1162 from patent US 6639063.
 25.1 GI:40168635

1.
 1.
 sified.
 ses 1 to 499)
 3, J.-B.D.M., Jobert, S. and Giordano, J.-Y.
 and encoded human proteins
 : US 6639063-A 1162 28-OCT-2003;
 Location/Qualifiers
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1.9%; Score 26; DB 6; Length 499;
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78 499 bp DNA PAT 18-SEP-2002
 78 encoded human protein.
 78.1 GI:23203896
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 ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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 ses 1 to 499)
 s, J.B.D.M., Jobert, S. and Giordano, J.B.
 d encoded human protein
 : JP 2002010789-A 1155 15-JAN-2002;
 CORP
 omo sapiens (human)
 P 2002010789-A/1155
 5-JAN-2002

PF 07-AUG-2000 JP 2000280989
 PR 05-AUG-1999 US 60/147499
 PI JEAN BAPTISTE DUMAS MILNE EDWARDS, SEVELIN JOBERT, JEAN
 GIORDANO
 PC C12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N
 C12N1/21,
 PC C12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12N5
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FEATURES
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 Matches 26; Conservative 0; Mismatches 0; Indels 0; G

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RESULT 55
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 ACCESSION AX885407
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 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 Dumas Milne Edwards, J.B., Duclert, A. and Giordano, J.Y.
 Expressed sequence tags and encoded human proteins
 Patent: EP 1033401-A 1270 06-SEP-2000;
 Genset (FR)

FEATURES
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 ACCESSION BD025017
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 KEYWORDS JP 2001269182-A/1263.

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 Meta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 es 1 to 516)
 is, J.B.D.M., Duclair, E. and Jordan, J.Y.
 ce tag and encoded human protein
 : JP 2001269182-A 1263 02-OCT-2001;
 :
 homo sapiens (human)
 JP 2001269182-A/1263
 02-OCT-2001
 24-FEB-2000 JP 2000118773
 26-FEB-1999 US 60/122487
 JEAN BAPTISTE DUMAS MILNE EDWARDS, EIMERIC DUCLAIR, JEAN YVES
 JORDAN
 12N15/09, C07K14/435, C07K16/18, C12N1/15, C12N1/19, C12N1/21, PC
 3/10,
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 934 522 bp DNA PAT 18-DEC-2003
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 ases 1 to 522)
 ds, J.-B.D.M., Jobert, S. and Giordano, J.-Y.
 and encoded human proteins
 t: US 6639063-A 3571 28-OCT-2003;
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 JP 2002010789-A/3564.
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 1 (bases 1 to 522)
 Edwards, J.B.D.M., Jobert, S. and Giordano, J.E.
 EST and encoded human protein
 Patent: JP 2002010789-A 3564 15-JAN-2002;
 GENSET CORP
 OS Homo sapiens (human)
 PN JP 2002010789-A/3564
 PD 15-JAN-2002
 PF 07-AUG-2000 JP 2000280989
 PR 05-AUG-1999 US 60/147499
 PI JEAN BAPTISTE DUMAS MILNE EDWARDS, SEVELIN JOBERT, JEA
 GIORDANO
 PC C12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12
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 ACCESSION AX381939
 VERSION AX381939.1 GI:19576761
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 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutel
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homc
 1
 Pyle, R.A., Xu, J. and Secrist, H.
 Amino acid sequences and methods for the therapy and diagnosis of
 cancer
 Patent: WO 0212280-A 877 14-FEB-2002;
 CORIXA CORPORATION (US)
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us-09-245-198a-3.oligo.rge

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22 22.1 GI:38328364

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ber,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
er,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
ul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
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enko,L., Maruina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
ton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
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ci,P., Prange,C., Raha,S.S., Loquellano,N.A., Peters,G.J.,
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an,K.J., Malek,J.A., Gunaratne,P.H., Richards,S.,
K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
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J., Helton,E., Kettner,M., Madan,A., Rodriguez,S.,
z,A., Whiting,M., Madan,A., Young,A.C., Shevchenko,Y.,
rd,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
field,Y.S., Krzywinska,M.I., Skalska,U., Smalusz,D.E.,
ch,A., Schein,J.E., Jones,S.J., and Marra,M.A.
tion and initial analysis of more than 15,000 full-length
and mouse cDNA sequences
Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
57
32
ses 1 to 534)
ber,R.
Submission
ted (13-NOV-2003) National Institutes of Health, Mammalian
ollection (MGC), Cancer Genomics Office, National Cancer
ute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
C Project URL: <http://mgc.nci.nih.gov>
t: MGC help desk
cgaps@mail.nih.gov
Procurement: ATCC
ibrary Preparation: CLONTECH Laboratories, Inc.
ibrary Arrayed by: The I.M.A.G.E. Consortium (LLNL)
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cgsc.bc.ca
Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield,
a Chan, Readman Chiu, Chris Fjell, Erin Garland, Ran Guin,
ia Hsiao, Martin Krzywinski, Reta Kutsche, Oliver Lee, Soo
e, Victor Ling, Carrie Mathewson, Candice McLeavy, Steven
Pawan Pandoh, Anna-Liisa Prabhu, Parvaneh Saeedi, Jacqueline
Duane Smalus, Michael Smith, Lorraine Spence, Jeff Stott,
Thorne, Miranada Tsai, Natasja van den Bosch, Jill Vardy,
Yang, Scott Zuyderduyn, Marco Marra.

distribution: MGC clone distribution information can be found
h the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
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Db 438 ATGTCATGTTAGACTTTGAAATTC 463

RESULT 61
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ACCESSION AR415931
VERSION AR415931.1 GI:40171041
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 538)
AUTHORS Edwards,J.-B.D.M., Jobert,S. and Giordano,J.-Y.
TITLE EST's and encoded human proteins
JOURNAL Patent: US 6639063-A 3568 28-OCT-2003;
FEATURES Location/Qualifiers
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Query Match 1.9%; Score 26; DB 6; Length 538;
Best Local Similarity 100.0%; Pred. No. 0.022;
Matches 26; Conservative 0; Mismatches 0; Indels 0; C

Oy 1 ATGTCATGTTAGACTTTGAAATTC 26
Db 451 ATGTCATGTTAGACTTTGAAATTC 476

RESULT 62

BD111484
LOCUS BD111484 538 bp DNA linear PAT 18
DEFINITION EST and encoded human protein.
ACCESSION BD111484
VERSION BD111484.1 GI:23206302
KEYWORDS JP 2002010789-A/3561.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 538)
AUTHORS Edwards,J.-B.D.M., Jobert,S. and Giordano,J.-Y.
TITLE EST and encoded human protein
JOURNAL Patent: JP 2002010789-A 3561 15-JAN-2002;
GENSET CORP

COMMENT

OS Homo sapiens (human)
PN JP 2002010789-A/3561
PD 15-JAN-2002
PF 07-AUG-2000 JP 2000280989
PR 05-AUG-1999 US 60/147499
PI JEAN BAPTISTE DUMAS MILNE EDWARDS, SEVELIN JOBERT, JEAN
GIORDANO
PC C12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N
C12N1/21,
PC C12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12NE
C12N15/00
CC EST and encoded human protein
FH Key Location/Qualifiers

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CDS      316..492.
          Location/Qualifiers
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          TCATTGTTAGACTTTGAAATTC 26
          |||||||
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          406 nce 1269 from Patent EPI033401. linear PAT 18-DEC-2003
          406
          406.1 GI:40041535
          sapiens (human)
          sapiens
          yota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          lia; Euthera; Primates; Catarrhini; Hominidae; Homo.

          Milne Edwards, J.B., Duclert, A. and Giordano, J.Y.
          ssed sequence tags and encoded human proteins
          t: EP 1033401-A 1269 06-SEP-2000;
          t (FR)

          Location/Qualifiers
            1..540
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              367..>540
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          1.9%; Score 26; DB 6; Length 540;
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          TCATTGTTAGACTTTGAAATTC 26
          |||||||
          TCATTGTTAGACTTTGAAATTC 527

          016 nce tag and encoded human protein. linear PAT 27-AUG-2002
          016
          016.1 GI:22566239
          01269182-A/1262.
          sapiens (human)
          sapiens
          yota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          lia; Euthera; Primates; Catarrhini; Hominidae; Homo.
          ases 1 to 540)
          ds, J.B.D.M., Duclair, E. and Jordan, J.Y.
          nce tag and encoded human protein
          t: JP 2001269182-A 1262 02-OCT-2001;
          T
          Homo sapiens (human)

```

```

PN JP 2001269182-A/1262
PD 02-OCT-2001
PF 24-FEB-2000 JP 2000118773
PR 26-FEB-1999 US 60/122487
PI JEAN BAPTISTE DUMAS MILNE EDWARDS, EIMERIC DUCLAIR, JE
PJ JORDAN
PC C12N15/09, C07K14/435, C07K16/18, C12N1/15, C12N1/19, C12
C12N5/10,
PC C12P21/02, C12P21/08, C12Q1/68//G06F17/30, C12N15/00, C1
G06F15/40
CC
FH Key Location/Qualifiers
FT CDS 367..540.

FEATURES
source
  Location/Qualifiers
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  |||||||
  Db . 502 ATGTCATTGTTAGACTTTGAAATTC 527

RESULT 65
BD224094 540 bp DNA linear PAT 1
LOCUS
DEFINITION Sequence characteristic to gene transcription controlled
          hypoxia.
ACCESSION BD224094
VERSION BD224094.1 GI:33033864
KEYWORDS JP 2002525081-A/9.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutel
          Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo
          1 (bases 1 to 540)
          Binat, P., Skalliter, R. and Feinstein, E.
          Sequence characteristic to gene transcription controlled
          TITLE Patent: JP 2002525081-A 9 13-AUG-2002;
          JOURNAL QUARK BIOTECH INC
          OS Homo sapiens (human)
          PN JP 2002525081-A/9
          PD 13-AUG-2002
          PF 27-AUG-1999 JP 2000571058
          PR 27-AUG-1998 US 60/098158, 05-MAY-1999 US 60/132
          PAZ EINAT, RAMI SKALITER, ELENA FEINSTEIN
          PC C12N15/09, A61K31/711, A61K45/00, A61K48/00, A61P9/00, A6
          A61P43/00,
          PC C12Q1/68, G01N33/53, G01N33/566, G01N33/58, G01N33/68, G0
          PC C12N15/00
          CC Sequence characteristic to gene transcription contro
          hypoxia
          FH Key Location/Qualifiers
          FT source 1..540
          FT /Organism="Homo sapiens (human)".

FEATURES
source
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ORIGIN
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  QY 1 ATGTCATTGTTAGACTTTGAAATTC 26

```

|||||
 CATTTAGACTTTGAAATTC 512

35 ce 3572 from patent US 6639063. linear PAT 18-DEC-2003

35 35.1 GI:40171045

n.

n.

sified.

ses 1 to 555)

S.J.-B.D.M., Jobert, S. and Giordano, J.-Y.

and encoded human proteins

: US 6639063-A 3572 28-OCT-2003;

Location/Qualifiers

1..555

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arity 1.9%; Score 26; DB 6; Length 555;

conservative 0; Mismatches 0; Indels 0; Gaps 0;

CATTTAGACTTTGAAATTC 26

|||||

CATTTAGACTTTGAAATTC 498

88 d encoded human protein. linear PAT 18-SEP-2002

88

88.1 GI:23206306

2010789-A/3565.

apiens (human)

apiens

ia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

ota; Eutheria; Primates; Catarrhini; Hominidae; Homo.

ses 1 to 555)

S.J.-B.D.M., Jobert, S. and Giordano, J.-Y.

d encoded human protein

: JP 2002010789-A 3565 15-JAN-2002;

CORP

omo sapiens (human)

P 2002010789-A/3565

5-JAN-2002

7-AUG-2000 JP 2000280989

5-AUG-1999 US 60/147499

EAN BAPTIST DUMAS MTINE EDWARDS, SEVELIN JOBERT, JEAN EVE

ANO

12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, PC

/21, 12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12N5/00, PC

/00

=a, g, c or t

Location/Qualifiers

338..514

IS

disc feature 541

Location/Qualifiers

1..555

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conservative 0; Mismatches 0; Indels 0; Gaps 0;

Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATGTCATTTAGACTTTGAAATTC 26

|||||

Db 473 ATGTCATTTAGACTTTGAAATTC 498

RESULT 68

BC042807

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Homo sapiens

REFERENCE

AUTHORS

TITLE

JOURNAL

Submitted (02-JAN-2003)

National Institutes of Health, Man

Gene Collection (MGC), Cancer Genomics Office, National Ce

Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892

USA

NIH-MGC Project URL: http://mgc.nci.nih.gov

Contact: MGC help desk

Email: cga@bmc.tmc.edu

Tissue Procurement: Miklos Palkovits, M.D., Ph.D.

cDNA Library Preparation: Michael J. Brownstein (NHGRI) &

Toshiyuki and Piero Carninci (RIKEN)

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Baylor College of Medicine Human Genome

Sequencing Center

Center code: BCM-HGSC

Web site: http://www.hgsc.bcm.tmc.edu/cdna/

Contact: amge@bcm.tmc.edu

Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Louis

Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanan

A.N., Gibbs, R.A.

Clone distribution: MGC clone distribution information car
 through the I.M.A.G.E. Consortium/LLNL at: http://image.ll
 Series: IRAK Plate: 91 Row: g Column: 13.

Location/Qualifiers

1..560

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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 0.022;

Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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|||||

Db 478 ATGTCATTTAGACTTTGAAATTC 503

RESULT 69

AR415932

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

Sequence 3569 from patent US 6639063.

AR415932.1 GI:40171042

Unknown.


```

4n.
ssified.
ases 1 to 595)
is,J.-B.D.M., Jobert,S. and Giordano,J.-Y.
and encoded human proteins
: US 6639063-A 3569 28-OCT-2003;
Location/Qualifiers
1. .595
/organism="unknown"
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Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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|||||
TCATTGTAGACTTTGAAATTC 538

485
nd encoded human protein.
485
485.1 GI:23206303
02010789-A/3562.
sapiens (human)
yota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
lia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ases 1 to 595)
ds,J.B.D.M., Jobert,S. and Giordano,J.E.
nd encoded human protein
t: JP 2002010789-A 3562 15-JAN-2002;
T CORP
Homo sapiens (human)
JP 2002010789-A/3562
15-JAN-2002
07-AUG-2000 JP 200280989
05-AUG-1999 US 60/147499
JEAN RAPUTIST DUMAS MILNE EDWARDS,SEVELIN JOBERT,JEAN EVE PI
DANO
C12N15/09,C12N15/09,C07K14/47,C07K16/18,C12N1/15,C12N1/19,PC
1/21
C12N5/10,C12P21/02,C12P21/08,C12Q1/68,C12N15/00,C12N5/00,PC
5/00
EST and encoded human protein
Key Location/Qualifiers
378. .554.
CDS Location/Qualifiers
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larity 100.0%; Pred.No.0.022;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

TCATTGTAGACTTTGAAATTC 26
|||||
TCATTGTAGACTTTGAAATTC 538

060
sapiens cdna FLJ26550 fis, clone LNF01586, highly similar to
aldolase (EC 2.2.1.2).
060

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VERSION AK130060.1 GI:34526798
KEYWORDS oligo capping; fis (full insert sequence) .
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Tashiro,H., Yamazaki,M., Watanabe,K., Kumagai,A., Itakura
Fukuzumi,Y., Fujimori,Y., Komiya,M., Suzuki,Y., Hata,H.
Nakagawa,K., Mizuno,S., Morinaga,M., Kawamura,M., Sugiyam
Irie,K., Otsuki,T., Sato,H., Nishikawa,T., Sugiyama,A.,
Kawakami,B., Nagai,K., Isogai,T. and Sugano,S.
NEDO human cDNA sequencing project
2 (bases 1 to 1822)
Unpublished
REFERENCE 2 (bases 1 to 1822)
AUTHORS Sugano,S. and Suzuki,Y.
TITLE Direct Submission
JOURNAL Submitted (31-JUL-2003) Sumio Sugano, Institute of Medica
University of Tokyo, Laboratory of Genome Structure, Huma
Center; Shirokane-dai, 4-6-1, Minato-ku, Tokyo 108-8639,
(E-mail:flcdna@ims.u-tokyo.ac.jp, Tel:81-3-5449-5286,
Fax:81-3-5449-5416)
COMMENT NEDO human cDNA sequencing project supported by Ministry
Economy, Trade and Industry of Japan; cDNA full insert se
Research Association for Biotechnology (RAB); cDNA librar
construction and 5'-end one pass sequencing: Institute of
Science, University of Tokyo, Laboratory of Genome Struct
Genome Center; 3'-end one pass sequencing: RAB; clone sel
full insert sequencing: RAB and Helix Research Institute.
FEATURES
source
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ORIGIN
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Best Local Similarity 100.0%; Pred.No.0.025;
Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 1 ATGTCATTGTAGACTTTGAAATTC 26
Db 1756 ATGTCATTGTAGACTTTGAAATTC 1781

RESULT 72
LOCUS HSM802687 2183 bp mRNA linear PRI 1
DEFINITION Homo sapiens mRNA; cDNA DKFZp762B195 (from clone DKFZp762
ACCESSION AL359585
VERSION AL359585.1 GI:8655645
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutel
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homc
1 (bases 1 to 2183)
Bloeker,H., Boecher,M., Brandt,P., Mewes,H.W., Weil,B. a
Wiemann,S.
Direct Submission
TITLE Submitted (15-JUN-2000) MIPS, Am Klopferspitz 18a, D-8215
JOURNAL Martinsried, GERMANY
COMMENT Clone from S. Wiemann, Molecular Genome Analysis, German
Research Center (DKFZ); Email s.wiemann@kfz-heidelberg.c
sequenced by GGF (National Research Centre for Biotechnol
Braunschweig/Germany) within the cDNA sequencing consorti
German Genome Project.
This clone (DKFZp762B195) is available at the RZPD in Ber

```

contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
-Charlottenburg, GERMANY; Email: clone@rzpd.de Further
info about the clone and the sequencing project is available
at: <http://www.mips.biochem.mpg.de/proj/cDNA/>.

Location/Qualifiers
1. .2183
/organism="Homo sapiens"
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DH10B; sites NotI + SalI"
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2140. .2145
2152

1.9%; Score 26; DB 9; Length 2183;
Identity 100.0%; Pred. No. 0.025;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ATTGTTAGACTTTGAAATTC 26
|||||
ATTGTTAGACTTTGAAATTC 2116

75 2650 bp mRNA linear PRI 09-SEP-2003
75 cDNA FLJ42181 f1s, clone THYMU2031368.
75.1 GI:34529902
capping; f1s (full insert sequence).
apiens (human)
apiens
ata; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

A., Takahashi-Fujii, A., Tanase, T., Imose, N., Takeuchi, K.,
A., Musashino, K., Yuuki, H., Hara, H., Sugiyama, T., Irie, R.,
T., Sato, H., Ota, T., Wakamatsu, A., Ishii, S., Yamamoto, J.,
I., Kawai-Hio, Y., Saito, K., Nishikawa, T., Kimura, K.,
ita, H., Matsuo, K., Nakamura, Y., Sekine, M., Kikuchi, H.,
C., Wagatsuma, M., Murakawa, K., Kanehori, K., Sugiyama, A.,
ni, B., Suzuki, Y., Sugano, S., Nagahara, K., Masuno, Y., Nagai, K.,
ngai, T.

man cDNA sequencing project

ished
ses 1 to 2650)

.T. and Yamamoto, J.

Submission

ed (15-JUL-2003) Takao Isegai, FLJ Project (HRI Team); 2-6-7
-Kamatori, Kisarazu, Chiba 292-0818, Japan
l:genomics@hri.co.jp, Tel:81-438-52-3975, Fax:81-438-52-3986)
man cDNA sequencing project supported by Ministry of
/ Trade and Industry of Japan; cDNA full insert sequencing:
th Association for Biotechnology (RAB); cDNA library
chnology: Helix Research Institute (HRI) (supported by Japan
chnology Center etc.); 5'- & 3'-end one pass sequencing: RAB,
id Biotechnology Center, National Institute of Technology and
ion; clone selection for full insert sequencing: HRI and
notation: HRI and RAB.

Location/Qualifiers

1. .2650
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/note="cloning vector: pME18SFL3"

ORIGIN

Query Match 1.9%; Score 26; DB 9; Length 2650;
Best Local Similarity 100.0%; Pred. No. 0.026;
Matches 26; Conservative 0; Mismatches 0; Indels 0; G

QY 1 ATGTCATTGTTAGACTTTGAAATTC 26
Db 2584 ATGTCATTGTTAGACTTTGAAATTC 2609

RESULT 74

HSDJ686N3

LOCUS

DEFINITION

HSDJ686N3 110293 bp DNA linear PRI 06
Human DNA sequence from clone RP4-686N3 on chromosome 20q1
Contains the 3' part of the gene for a novel ATP dependent
helicase (contains conserved C-terminal helicase domains a
DEAD/DEAH boxes), the KIAA1404 gene, a putative novel gene
STSs, GSSs and two CpG islands, complete sequence.

AL049766

AL049766.14 GI:5763746

HTG; CpG island; DEAD box; DEAH box; KIAA1404; RNA helicase

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 110293)

Corby, N.

Direct Submission

Submitted (01-MAR-2001) Sanger Centre, Hinxton, Cambridges

CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

requests: clonerequest@sanger.ac.uk

On Aug 24, 1999 this sequence version replaced gi:5730221.

During sequence assembly data is compared from overlapping

Where differences are found these are annotated as variati

together with a note of the overlapping clone name. Note t

variation annotation may not be found in the sequence subm

corresponding to the overlapping clone, as we submit sequ

only a small overlap as described above.

The following abbreviations are used to associate primary

numbers given in the feature table with their source datab

Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Info

on the WORMPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This se

the entire insert of clone RP4-686N3 this sequence was fin

follows unless otherwise noted: all regions were either

double-stranded or sequenced with an alternate chemistry o

by high quality data (i.e., phred quality >= 30); an attem

made to resolve all sequencing problems, such as compressi

repeats; all regions were covered by at least one plasmid

or more than one M13 subclone; and the assembly was confir

restriction digest. This sequence was generated from part

bacterial clone contigs of human chromosome 20, constructe

Sanger Centre Chromosome 20 Mapping Group. Further inform

be found at <http://www.sanger.ac.uk/HGP/Chr20>

RP4-686N3 is from the library RPCI-4 constructed by the gr

Pieter de Jong. For further details see

<http://www.chori.org/bacpac/home.htm>

VECTOR: pCYPAC2.

Location/Qualifiers

1. .110293

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/note="AluY repeat: matches 2. .62 of consensus"

207. .21327

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5965. .6138,7449. .7599,10556. .10655,10818. .10959,12185. .12338,13400. .13481,13583. .13760,16199. .16305,16486. .16588,19143. .19237,19333. .19456,19844. .19931,20879. .20953,21066. .21327)
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849. .1146
/notes="AluSx repeat: matches 1. .298 of consensus"
1147. .1423
/notes="AluJo repeat: matches 4. .282 of consensus"
1605. .1912
/notes="AluSx repeat: matches 1. .307 of consensus"
complement(1913. .2134)
/notes="match: GSS: Em:AQ349240"
1952. .2099
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2473. .2589
/notes="FLAM_C repeat: matches 2. .125 of consensus"
2617. .2703
/notes="L2 repeat: matches 2051. .2135 of consensus"
2704. .3004
/notes="AluY repeat: matches 1. .301 of consensus"
3005. .3572
/notes="L2 repeat: matches 2135. .2708 of consensus"
3908. .4051
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4108. .4411
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AluYa5 repeat: matches 1. .304 of consensus"
4441. .4610
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5392. .5685
/notes="AluSg repeat: matches 1. .301 of consensus"

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5690. .5809
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AluSg/x repeat: matches 14. .133 of consensus"
6281. .6579
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6896. .7199
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/note="MIR repeat: matches 48. .174 of consensus
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8870. .9009
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9126. .9436
/note="AluJo repeat: matches 1. .308 of consensu
9437. .9492
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9598. .9731
/note="AluSp repeat: matches 1. .127 of consensu
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/note="L1MC4 repeat: matches 7624. .7814 of cons
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/note="MIR repeat: matches 178. .250 of consensu
12997. .13292
/note="AluY repeat: matches 1. .296 of consensu
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complement(13759. .14174)
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/gene="dJ686N3.1"
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/gene="dJ686N3.1"
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14175. .14282
/note="AluJ/FLAM repeat: matches 2. .87 of conse
AluJ/FLAM repeat: matches 2. .87 of consensus"
14304. .14596
/note="AluJo repeat: matches 3. .300 of consensu
14744. .15145
/note="L2 repeat: matches 2251. .2749 of consens
15261. .15558
/note="AluSc repeat: matches 9. .306 of consensu
15560. .15717
/note="AluJo repeat: matches 145. .293 of consen
15718. .16016
/note="AluSx repeat: matches 1. .297 of consensu

Query Match 1.9%; Score 26; DB 9; Length 110293;
Best Local Similarity 100.0%; Pred. No. 0.035;
Matches 26; Conservative 0; Mismatches 0; Indels 0;

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06:25:14 2004

us-09-245-198a-3.oligo.rge

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arity 100.0%; Pred.No. 0.036;
conservative 0; Mismatches 0; Indels 0; Gaps 0;

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April 8, 2004, 22:35:57
acs

16:25:16 2004

us-09-245-198a-3.oligo.rst

GenCore version 5.1.6
copyright (c) 1993 - 2004 Compugen Ltd.

c search, using sw model

il 8, 2004, 20:38:11 ; Search time 3960 seconds
(without alignments)
10353.731 Million cell updates/sec

-09-245-198a-3
3
tgtcattgtgacttga.....gacaaaattgtgataatgg 1373

GO_NUC

op 60.0 , Gapext 60.0

13289 seqs, 14931090276 residues

s satisfying chosen parameters: 55026578

th: 0

th: 2000000000

sting first 100 summaries

T:*

em_estba:*

em_esthum:*

em_estin:*

em_estnu:*

em_estov:*

em_estpl:*

em_estro:*

em_htc:*

gb_est1:*

gb_est2:*

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em_gss_fun:*

em_gss_mam:*

em_gss_mus:*

em_gss_pro:*

em_gss_rod:*

em_gss_pbg:*

em_gss_vrl:*

gb_gss1:*

gb_gss2:*

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
ad by analysis of the total score distribution.

SUMMARIES

ry	ch	Length	DB	ID	Description
1.1	776	13	BX090012	BX090012	
1.8	777	12	BI819200	BI819200	603034614
1.6	1071	12	BM921213	BM921213	AGENCOURT
1.1	963	13	BQ671259	BQ671259	AGENCOURT

CA396679 C
CB141389 K
BI596681 6
BI870393 6
BI871711 6
CA413067 U
BI966060 I
CF126539 U
BU951915 I
CB529199 U
BQ707185 A
BQ884231 A
CF126932 U
BI766766 6
AI422796 tf
BM971606 U
BU631264 U
BI824443 6
CB998034 A
BM128059 I
BQ674188 A
BM509016 I
BM562622 U
BM658822 7
AW195034 X
BF439993 R
AW204512 U
AW291620 U
AI913541 wa
BI966255 I
AQ100365 H
BM925491 A
BM505649 I
AI695776 wb
AI221985 qg
AI291866 qm
BG110063 6
AI202121 q1
BM703512 U
BU729427 U
BM688946 U
BM686319 6
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AI760777 w1
BF940141 n
BG054878 R
AW081731 xb
AW661741 h
AI863563 w1
AI682487 w1
BI906850 6
BG054914 n
AI669243 wc
BI908274 6
AI091441 ow
BF222608 7
CO0994 HUMK
CF994566 AC
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BE501197 7
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AI081661 ou
N35070 YL
BI965174 I

8.7	639	10	BB643326
8.7	692	13	B7489662
8.7	1187	12	B0053284
7.7	2237	11	AK044387
7.4	1926	12	BM906056
7.4	2854	12	BM503779
7.1	259	12	BM1728317
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6.9	837	14	AW911845
6.8	105	10	BF041509
6.6	538	10	BF821434
6.1	140	9	AL582027
5.9	576	9	AL535912
5.3	278	10	AF417023
5.3	404	10	BF044430
5.3	542	10	BF041509
5.2	257	14	R55295
5.2	257	14	R55295
5.2	784	12	BG686384
4.2	571	10	BF073981
3.8	584	10	AW911574
3.6	426	10	AW236926
3.6	561	10	AW763237
3.4	237	29	CG616340
3.4	269	29	CG564287

ALIGNMENTS

12 Soares breast 2NbHbSt Homo sapiens cDNA clone
776 bp mRNA linear EST 23-JAN-2003
98E15243 ; IMAGE:154742, mRNA sequence.

12.1 GI:27821952

apiens (human)

apens (unknown)
 Crata; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinoptera; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 ses 1 to 776)
 L., Heil O., Hennig S., Neubert P., Partsch E., Peters M.,
 f U., Schneider D. and Korn B.
 Unigeneset - RZPD3
 ished (2003)
 t: Ina Rolfs
 eutsches Ressourcenzentrum fuer Genomforschung GmbH
 enhelmer Feld 580, D-69120 Heidelberg, Germany
 INAGP998E15243.

B; I.M.A.G.E. cDNA Clone Collection;

UnigeneSet - RZPD3 (RZPDLIB No.972)

[/www.rzpd.de/CloneCards/cgi-](http://www.rzpd.de/CloneCards/cgi-)

owLib.pl.cgi/response?LibNo=972 Cont

deutsches Ressourcenzentrum fuer Genomforschung (DZG), Leipzig

rweg 6, D-14059 Berlin, Germany

49 30 32639 101

49 30 32639 111

pd.de

None is available royalty-free from

t RZPD (clone@rzpd.de) for further in-

Primer sequence: TTTACACAGGAAACAGCTT

Location/Qualifiers

1. .776

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/organism="Homo sapiens"
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/mol_type="mRNA"
/dh_mtf="4.7777777777777777"

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/db_xref="taxon:9606"
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/clone="IMAGp998E15243" ; IMAGE:15
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/sex="Female"  
/dev et age="adul" + "
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/lab_host="DHIOB (ampicillin res)
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/clone_lib="Soares breast 2NbHBst"
/note="Organ: breast: Vector: nT7"
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/note="Organ: breast; Vector: p1"/
modified polymer: site: Not
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modified polylinker; site_1: Not

strand cDNA was primed with a Not I - oligo (dT).
TGTTCACATCTGAGTCGGCCGCCCGCCCCCTTTTCTTTTTT
Double stranded cDNA was ligated to Eco RI adapt
(pharmacia), digested with Not I and cloned into
and Eco RI sites of a modified p7r3 vector (Phar
Library went through one round of normalization.
230. Library constructed by Bento Soares and M.Fa
Bonaldi."

ORIGIN

Query Match	52.1%	Score 716;	DB 13;	Length 776;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 716;	Conservative 0;	Mismatches 0;	Indels 0;	
QY	600	TCAGTGTGCACATTGATCAGGGGAAGCGTGTACCTGAGAGCTGGAGCTGCTGGTGG		
Db	1	TCAGTGTGCACATTGATCAGGGGAAGCGTGTACCTGAGAGCTGGAGCTGCTGGTGG		
QY	660	TGTGTCGGCCCTGCGCTGCGCTCGAGGAAATTTCTCAGCCACTGCGGGCCAGTTCCCTCC		
Db	61	TGTGTCGGCCCTGCGCTGCGCTCGAGGAAATTTCTCAGCCACTGCGGGCCAGTTCCCTCC		
QY	720	CCAGTCTGCGCTCTGCGCAGGTGCTGGGCTGTGGGCTGTGGCCCTGCGGCCAGGGTCTCTCC		
Db	121	CCAGTCTGCGCTCTGCGCAGGTGCTGGGCTGTGGGCTGTGGCCCTGCGGCCAGGGTCTCTCC		
QY	780	GATCGGACACCTCCCTGGGGCCCATCTCAAGGCTGCCGCCCTTCTCCTCACTACTTCC		
Db	181	GATCGGACACCTCCCTGGGGCCCATCTCAAGGCTGCCGCCCTTCTCCTCACTACTTCC		
QY	840	CTTCCAGGTTCACTGAGGGGCCCTGTGTTCTCCCAAGTGTGCCCAGGCTGCCGGCT		
Db	241	CTTCCAGGTTCACTGAGGGGCCCTGTGTTCTCCCAAGTGTGCCCAGGCTGCCGGCT		
QY	900	TCGACAGCTCTGTGGGCAACCGGTGCCCTCTGCCCAAGTGTGCCCAGGCTGCCGGCT		
Db	301	TCGACAGCTCTGTGGGCAACCGGTGCCCTCTGCCCAAGTGTGCCCAGGCTGCCGGCT		
QY	960	ACCTGCCCTCCCTCAGAGGCTGCTGGGCTGTTCAGCTGTTTCCATCCCACT		
Db	361	ACCTGCCCTCCCTCAGAGGCTGCTGGGCTGTTCAGCTGTTTCCATCCCACT		
QY	1020	TACAGTATTCCCACTCTTATCTTACAACTTCCCCCAAGGCTGCCCTCCTCCTCCT		
Db	421	TACAGTATTCCCACTCTTATCTTACAACTTCCCCCAAGGCTGCCCTCCTCCTCCT		
QY	1080	CCCAATCCCTGACCTTTGAGGCCCCCAGTGTGATCTGACTGCCCTGGGCCACAG		
Db	481	CCCAATCCCTGACCTTTGAGGCCCCCAGTGTGATCTGACTGCCCTGGGCCACAG		
QY	1140	CCAGGCAATTGTCTACTCTCTGTGGGCAAGGATGGGTCAGAGAGACCCCACT		
Db	541	CCAGGCAATTGTCTACTCTCTGTGGGCAAGGATGGGTCAGAGAGACCCCACT		
QY	1200	GGCACTAAGAGGGGCTGGACCTTGGCGGCAAGGACCAAGAGACTGGGGCTTAGGCC		
Db	601	GGCACTAAGAGGGGCTGGACCTTGGCGGCAAGGACCAAGAGACTGGGGCTTAGGCC		
QY	1260	GTTCCCAAATGTGAGGGGGAGAGAAACAGCAAGCTCTCTCCTTGAGAAATTCCTCT		
Db	661	GTTCCCAAATGTGAGGGGGAGAGAAACAGCAAGCTCTCTCCTTGAGAAATTCCTCT		

RESULT 2

AB030052	BI819200	777 bp	linear	EST 04
LOCUS	60303461.f1	NIH MGC_115	Homo sapiens cDNA clone IMAGE:5175	
DEFINITION	mRNA sequence.			
ACCESSION	BI819200			
VERSION	BI819200			
KEYWORDS	BI819200.1	GI:15930750		
SOURCE	EST.			
ORGANISM	Homo sapiens (human)			
	Homo sapiens			

ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 777

http://mhc.nci.nih.gov/
l. Institutes of Health, Mammalian Gene Collection (MGC)
shed (1999)

: Robert Strausberg, Ph.D.

cgapbs-remail.nih.gov

Procurement: Life Technologies, Inc.

Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

quencing by: Incyte Genomics, Inc.

distribution: MGC clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:

image.llnl.gov

LLAM1437 row: 1 column: 03

ality sequence stop: 759.

Location/Qualifiers

1. .777

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:5175698"

/lab_host="DH10B"

/clone_lib="NIH MGC 115"

/note="Organ: pooled brain, lung, testis; Vector:

pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA

source anonymous pool of 6 male brains, age range 23-27; 1

male lung, age 27, and 1 male testis, age 69. Library is

oligo-dr primed and directionally cloned (EcoRV site is

destroyed upon cloning). Average insert size 1.8 kb,

insert size range 1-3 kb. Library is normalized and

enriched for full-length clones and was constructed by C.

Gruber (Invitrogen). Research Genetics tracking code

021. Note: this is a NIH_MGC Library."

45.8%; Score 629; DB 12; Length 777;

arity 99.7%; Pred. No. 3.1e-309;

conservative 0; Mismatches 2; Indels 0; Gaps 0;

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CTTTCTGAAACCGACTAGTTGGGCTCGCAGAGTGACCTTAAGCGCGGAAC 132

|||||

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|||||

AGGAGAGTGTGGACGGACAGTGAAGTGGCTGGGAGGAGGAGGAGGAGGAGG 252

|||||

CTCTGCGCTACAAACCGCAGATCGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 587

|||||

CTCTGCGCTACAAACCGCAGATCGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 312

|||||

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|||||

TGTAAGTGTGAGTGTGAGTGTGAGTGTGAGTGTGAGTGTGAGTGTGAGTGTG 372

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TGGATGTGTGTGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG 432

|||||

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|||||

TCGGGCCCCAGCTCCGCTCTCGCCAGGTGCTGGGCTGTGGGCTGTGGGCTGT 492

768 GTCTCTCCCTCGGATCCGACCCCTCCCTCGGCGCCCATCTCAAGGCTGCCCTTCTCT
|||||

493 GTCTCTCCCTCGGATCCGACCCCTCCCTCGGCGCCCATCTCAAGGCTGCCCTTCTCT
|||||

828 CTACTTCGAGCTCTTCCAGGTTCACTGAGGGGCCCTGTCTCCCCACAGTGTCTCCCP
|||||

553 CTACTTCGAGCTCTTCCAGGTTCACTGAGGGGCCCTGTCTCCCCACAGTGTCTCCCP
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888 TGCAGGCTCCCTCGACAGCTCTCTGGGACACCGGTCCTCTGCCCTTCTGCCCTCAGC
|||||

613 TGCAGGCTCCCTCGACAGCTCTCTGGGACACCGGTCCTCTGCCCTTCTGCCCTCAGC
|||||

948 TCTTTGTCTCCAGACCTGCCCTCTCTAGAGGCTGCTGGGCTCTTTCACGTGTT
|||||

673 TCTTTGTCTCCAGACCTGCCCTCTCTAGAGGCTGCTGGGCTCTTTCACGTGTT
|||||

1008 ATCCACATAA 1018

|||||

733 ATCCACATAA 743

|||||

RESULT 3

BM921213

LOCUS

DEFINITION

5', mRNA sequence.

BM921213

BM921213.1 GI:19371592

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutelec

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 1071)

NIH-MGC http://mhc.nci.nih.gov/

National Institutes of Health, Mammalian Gene Collection (M

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-remail.nih.gov

Tissue Procurement: Life Technologies, Inc.

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Agencourt Bioscience Corporation

Clone distribution: MGC clone distribution information can

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLAM12786 row: p column: 02

High quality sequence stop: 656.

Location/Qualifiers

1. .1071

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:5752561"

/lab_host="DH10B"

/clone_lib="NIH MGC 115"

/note="Organ: pooled brain, lung, testis; Vector:

pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroye

source anonymous pool of 6 male brains, age range

male lung, age 27; and 1 male testis, age 69. Lib

oligo-dr primed and directionally cloned (EcoRV si

destroyed upon cloning). Average insert size 1.8

insert size range 1-3 kb. Library is normalized an

enriched for full-length clones and was construct

Gruber (Invitrogen). Research Genetics tracking

021. Note: this is a NIH_MGC Library."

ORIGIN

Query Match

Best Local Similarity 44.6%; Score 613; DB 12; Length 1071;

Matches 713; Conservative 0; Mismatches 2; Indels 0; Ga

412 GCTCGAAGAGCGATCGCAGCCCATTTATGAAGTTTCATCCAGCAGCTGGACGACGGA

.y1 Human Retinal pigment epithelium/choroid cDNA
 malized, unamplified): cs Homo sapiens cDNA clone cs80h07
 A sequence.

9.1 GI:24733297

piens (human)

ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 es 1 to 567)
 G., Bernstein,S.L., Wyatt,M.K., Farris,R.N., Behal,A.,
 n,J.W., Bouffard,G., Smith,D. and Peterson,K.
 ed sequence tag analysis of human RPE/choroid for the
 Project: Over 6000 non-redundant transcripts, novel genes
 ice variants
 s. 8 (4), 205-220 (2002)

0

0

on Molecular Structure and Function

1 Eye Institute

NIH, Bethesda, MD 20892-2740, USA

1 402 3452

1 496 0078

graeme@helix.nih.gov

80 row: h column: 07

mer: M3RPI reverse primer (ABI).

Location/Qualifiers

1. .567

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="cs80h07"

/tissue_type="RPE/choroid"

/dev_stage="Adult"

/lab_host="EMDH10B"

(Un-normalized, unamplified): cs"
 /note="Organ: Eye; Vector: pCMVSPORT6; Two different donor
 eyes (75-80 years old) yielded approximately 600 mg of
 dissected RPE/choroid tissue. This in turn yielded 340 ug
 of total RNA and 7 ug of mRNA. A directionally cloned cDNA
 library in the pCMVSPORT6 vector was constructed at Life
 Technologies (Rockville, MD; now part of Invitrogen Corp),
 essentially following the protocols of the SuperScript
 Plasmid System (Invitrogen Corp).
 <http://www.invitrogen.com/>". The library code
 designation was cs. For this library, cDNA inserts were
 cloned into the NotI/MluI sites of the vector. EST
 analysis was performed on the unamplified library at the
 NIH Intramural Sequencing Center (NISC)."

41.3%; Score 567; DB 14; Length 567;
 rity 100.0%; Pred. No. 1.3e-277;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;

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|||||

3TCCCGGATGGGGGGGGGTGAGGACAGGACAGCCCCCGCCCGATGCGCGC 60

|||||

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|||||

GAGCCAGAGCGGAGGGGGCGCCGGGGGAGCGGGACCGCCCTGCTGTCGCG 120

|||||

GCTGGGCTGCGGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTG 234

|||||

GCTGGGCTGCGGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTG 180

|||||

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|||||

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 |||||
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 QY 355 TTCCTGAACCGACTAGTTCGGCCTCGCAGAACTGCACTTAAAGCGCGGAAAAACACG
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 QY 415 CGAAGAGCGATCGGAGCCCATTTATGAAGTTCATCCAGCAGCTGGACGACGAGGCGC
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 Db 361 CGAAGAGCGATCGGAGCCCATTTATGAAGTTCATCCAGCAGCTGGACGAGGAGCGC
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 Db 541 TACTGTCAAGTGCACCTTTGATGAGGGG 567
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RESULT 6

LOCUS

CB141389

DEFINITION

K-EST0194999 L15CKK1 Homo sapiens cDNA clone L15CKK1-30-E06

mRNA sequence.

ACCESSION

CB141389

VERSION

CB141389.1 GI:28116436

KEYWORDS

EST.

SOURCE

Homo sapiens (human)

ORGANISM

Homo sapiens

REFERENCE

1 (bases 1 to 545)

AUTHORS

Kim,N.S., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., K

Kim,K.J., Cheong,J.E., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S

Kim,Y.S.

21C Frontier Korean EST Project 2001

Unpublished (2002)

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Fax: +82-42-860-4409

Email: yongsung@mail.kr.ibm.re.kr

Plate: 30 row: E column: 06

High quality sequence stop: 545.

Location/Qualifiers

1. .545

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/lab_host="Top10P"

/clone_lib="L15CKK1"

/note="Organ: Liver; Vector: pcNS-D2; Site: 1; EcoR

Site 2: NotI; The poly (A) + RNA was dephosphorylat

bacterial alkaline phosphatase (BAP) and then deca

with tabacco acid pyrophosphatase (TAP). The decap

intact mRNA was ligated with DNA-RNA linker includ

EcoRI site by treatment of T4 RNA ligase and the f

strand cDNA was synthesized from oligo dt-selected

priming with dt-tailed vector. The dt-tailed vecto

adjusted to have about 60nt. The cDNA vector was

circularized with E. coli DNA ligase after digesti

EcoRI which site is also included in vector. An RNA strand converted to a DNA strand by Okayama-Berg method. The obtained cDNA vectors were used for transformation of competent cells *E. coli* Top10[®] by electroporation method. The cDNA libraries constructed by this method are full-length enriched cDNA library."

39.7%; Score 545; DB 14; Length 545;
arity 100.0%; Pred. No. 2.1e-266;
conservative 0; Mismatches 0; Indels 0; Gaps 0;

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T 853
T 545

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sequence.

81.1 GI:15489620

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apiens
apiens
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Euthera; Primates; Catarrhini; Hominidae; Homo.
uses 1 to 828)
http://mgc.nci.nih.gov/.
al Institutes of Health, Mammalian Gene Collection (MGC)
ished (1999)
t: Robert Strausberg, Ph.D.
cgaps@remail.nih.gov
Procurement: Miklos Palkovits, M.D., Ph.D.
Library Preparation: Michael J. Brownstein
(NHGRI), Shiraki

Toshiyuki and Piero Carninci (RIKEN)
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information c2
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>

Plate: LLAM11722 row: k column: 13
High quality sequence stop: 776.

[illegible]

Conservativity	100.0%	Pred. No. 2.1e-266;	0;	Mismatches	0;	Indels	0;	Gaps	0;
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CA	TATGAAGTTCATCCACGACCTTGACAGACGCGAGCGAGCAGTGTGACGG	180							
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ORIGIN

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Best Local Similarity 99.7%; Pred. No. 1.4e-252;
Matches 618; Conservative 0; Mismatches 2; Indels 0; C
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Db 103 TCGGCCTCGCAGAAAGTGCACCTAAAGGCCGGAACAACCGGGCTCGAAGAGCGATCC

ov 432 CCATTATGAAGTTCATCCACGACTTGGACAGGACGGAGCGCGAGCGAGCTTGGACG

432 CCATTATGAGATTCATCCACGACCTGGACACGAGCGCAGGCAGGCTGGAC
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QY 552 GATCGGGGAGTTTATAGTCACCCGGGCTGGGCTCTACTACCTGTACTGTCAAGGTG

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Ov 612 TGATGAGGGGAAGGCTGTCTACCTGAAGCTGGACCTGCTGGTGGATGGTGTGCTG

QY
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Db
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QY 672 GGGTGCCTGGAGGAATTCTCAGCCACTGCGGCCAGTTCCTTCGGGGCCCCAGCTCC
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db 403 GGGTGCCTGGAGGAATTCTCAGCCACTGCGGCCAGTTCCTTCGGGGCCCCAGCTCC
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Db 583 CTGAGGGGGCCCTGGTCTCTCCCCCGAGTCGTCCCAGGCTGCCGGGTCCCCCTCGACAG

S., T., Jackson, Y. and Bowers, Y.
ne Pancreas Consortium
shed (2000)
: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
ne Pancreas Consortium
University, Howard Hughes Medical Institute
Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
8
7-495-1812
7-495-8557
dmelton@bichp.harvard.edu
was constructed by Dr. Douglas Melton DNA sequencing by:
ton University Genome Sequencing Center For information on
ng a clone please contact: Juliana Brown
fas.harvard.edu) This sequence now available from the IMAGE
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made by oligo-dT priming. Size-selected by column
fractionation; average insert size 1.08 kb. Library was
amplified once on solid support and plasmid DNA from
library was prepared. The library DNA was normalized by
method #4 from Bonaldo, Lennon, and Soares 1996 Genome
Research 6:791-806; 0.5 microgram single-stranded library
plasmid DNA was mixed with 5 micrograms PCR product
representing library inserts and hybridized to an EcoT of
20. Single-stranded (unhybridized) plasmids were isolated
by hydroxyapatite chromatography and used to make this
library."

35.8%; Score 491; DB 12; Length 609;
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REFERENCE 1 (bases 1 to 666)
AUTHORS Bonaldo, M.F., Lennon, G. and Soares, M.B.
TITLE Normalization and subtraction: two approaches to facilitate
discovery
JOURNAL Genome Res. 6 (9), 791-806 (1996)
MEDLINE 97044477
PUBMED 8889548
COMMENT Contact: Soares, MB
Coordinated Laboratory for Computational Genomics
University of Iowa
375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: bento-soares@uiowa.edu
Tissue Procurement: Mary Hendrix
cDNA Library preparation: Dr. M. Bento Soares, University
cDNA Library Arrayed by: Dr. M. Bento Soares, University
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Distribution information can be found
http://genome.uiowa.edu/distribution/humanfl.html
Seq primer: pyX-5.
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Site 2: Not I; The library was constructed accordi
Bonaldo, Lennon and Soares, Genome Research, 6:791
1996. Denatured RNA was size fractionated on a 1%
gel. First strand cDNA synthesis was primed with c
primer containing a Not I site. Double strand cDNA
size selected according to mRNA size fraction, lig
with EcoR I adaptor, digested with Not I and then
directionally into pyX-Asc vector. The library tag
sequence located between the Not I site and the po
is GATAAGGCCA. Tissue was provided by Mary Hendrix

Query Match 33.9%; Score 465; DB 14; Length 666;
Best Local Similarity 99.6%; Pred. No. 1.5e-225;
Matches 565; Conservative 0; Mismatches 2; Indels 0; G
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 ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 D., Brown, J., Kenty, G., Permut, A., Lee, C., Kaestner, K.,
 ka, I., Scarce, M., Brestelli, J., Gradwohl, G., Clifton, S.,
 L., Marra, M., Pape, D., Wylie, T., Martin, J., Bliscain, A.,
 t, A., Theising, B., Ratter, E., Ronko, I., Bennett, J.,
 as, M., Gibbons, M., McCann, R., Cole, R., Tsagareishvili, R.,
 ms, T., Jackson, Y., and Bowers, Y.
 ine Pancreas Consortium
 ished (2000)
 t: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
 ine Pancreas Consortium
 d University, Howard Hughes Medical Institute
 of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
 38
 17-495-1812
 17-495-8557
 dmelton@biohp.harvard.edu
 y was constructed by Dr. Hiroshi Inoue DNA sequencing by:
 ation University Genome Sequencing Center For information on
 ing a clone please contact: Dr. Hiroshi Inoue
 ce@im.wustl.edu
 1mer: -400P from Gibco

High quality sequence stop: 451.
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 Size-selected on agarose gel. Average insert size
 XhoI site was destroyed after directional cloning
 Amplified once. Contact information: Hiroshi Inoue
 Metabolism Div. (Alan Permut Lab), Washington Ur
 School of Medicine, Box 8127, 560 South Euclid Av
 Louis, MO 63110, E-mail: hinoue@imgate.wustl.edu,
 314-362-1916, Fax: 314-747-2692."

ORIGIN
 Query Match 33.6%; Score 462; DB 13; Length 474;
 Best Local Similarity 100.0%; Pred. No. 4.8e-224;
 Matches 462; Conservative 0; Mismatches 0; Indels 0; G

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 Db 294 ACCCTTTGAGGCCCCCAGTGTCTGACTCCCCCTGGCCACAGACCCCAAGGGC
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 Db 234 GTTCACTGTACTCTGTGGGCAAGGATGGTCCAGAGACCCCACTTCAGGCACTAT
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 QY 1272 GAGGGCGAGAAACAAGACAGCTCTCCCTTGAGAAATTCCTGTGGATTTTAA
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RESULT 14
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 VERSION CB529199.1 GI:29389647
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 SOURCE Homo sapiens (human)
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 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 569)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (

ane Index
shed (1997)
Robert Strausberg, Ph.D.
cgapbs-remail.nih.gov
Procurement: Dr. Gary W. Hunninghake, U of I
Library preparation: Dr. M. Bento Soares, University of Iowa
Library Arrayed by: Dr. M. Bento Soares, University of Iowa
Sequencing by: Dr. M. Bento Soares, University of Iowa
Distribution: Distribution information can be found at
genome.uiowa.edu/distribution/cgap.html
Following repetitive elements were found in this cDNA
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challenged with different treatments. The library was
subtracted according to Bonaldo, Lennon and Soares, Genome
Research, 6:791-806, 1996. The tissue was provided by Dr.
Gary W. Hunninghake of the University of Iowa.
TAG TISSUE=Human Lung Aveolar Macrophage
TAG LIB=UI-H-FT2
TAG_SEQ=GGCCATGCCG"

32.7%; Score 449; DB 14; Length 569;
rity 99.8%; Pred. No. 2.1e-217;
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QY 1354 GACAAATGTTGATAAATGG 1373
Db 47 GACAAATGTTGATAAATGG 28
RESULT 15
BQ707185
LOCUS
DEFINITION BQ707185 948 bp mRNA linear EST 16-
AGENCOURT 8353983 NIH_MGC_113 Homo sapiens cDNA clone IMAGE
5', mRNA sequence.
ACCESSION BQ707185
VERSION BQ707185.1 GI:21846084
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleo
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 948)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (M
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Dr. Mark Watson
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (ILNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can
found through the I.M.A.G.E. Consortium/ILNL at:
http://image.llnl.gov
Plate: LLCM2466 row: n column: 17
High quality sequence start: 24
High quality sequence stop: 550.
Location/Qualifiers
1..948
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6278608"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC_113"
/note="Organ: spleen; Vector: pOTB7; Site 1: XhoI;
EcoRI; cDNA made by oligo-dT priming. Directional
into EcoRI/XhoI sites using the following 5' adapt
GGCAGCAG(G). Library constructed by Ling Hong in
laboratory of Gerald M. Rubin (University of Cali
Berkeley) using ZAP-cDNA synthesis kit (Stratagene
Superscript II RT (Life Technologies)). Note: this
NIH_MGC Library."

ORIGIN

Query Match 31.8%; Score 437; DB 13; Length 948;
Best Local Similarity 99.8%; Pred. No. 3.2e-211;
Matches 487; Conservative 0; Mismatches 1; Indels 0; Ga
QY 475 GCAGTGTGACGCGGACGAGTGGCTGGGAGGAGCCAGAAATCAACAGCTCCAGC
Db 167 GCAGTGTGACGCGGACGAGTGGCTGGGAGGAGCCAGAAATCAACAGCTCCAGC
QY 535 CTGCGCTACACCGCCAGATCGGGAGTTTATAGTCAACCGGGCTGGGCTCTACTAC
Db 227 CTGCGCTACACCGCCAGATCGGGAGTTTATAGTCAACCGGGCTGGGCTCTACTAC
QY 595 TACTTCTCAGGTGCATTTGATGAGGGAGGAGCTGTCTACTCTGAGCTGAGCTTGTCTG
Db 287 TACTTCTCAGGTGCATTTGATGAGGGAGGAGCTGTCTACTCTGAGCTGAGCTTGTCTG
QY 655 GATGTGTGTGCTGGCCCTCGCTCGCTGAGGAAATTTCTCAGCCACTGCGGCCAGTTTCC
Db 347 GATGTGTGTGCTGGCCCTCGCTCGCTGAGGAAATTTCTCAGCCACTGCGGCCAGTTTCC
QY 715 GGGCCCCAGCTCGCCCTCTGCCAGGTGTCTGGGCTGTGGCCCTCGGCCCGAGGTCC

CCAGCTCCGCTCTGCGAGGTGTCTGGGCTGTGGCCCTTGGCCCTCGCGCCAGGGTCTCTCC 466
 GGATCCGACACCTCCCTGGGCCCATCTCAAGGCTGCCCTTCTCACTTCTTCTCACTTCTT 834
 GGATCCGACACCTCCCTGGGCCCATCTCAAGGCTGCCCTTCTCACTTCTTCTCACTTCTT 526
 TCTTCCAGGTTCACTGAGGGGCTCTGTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTT 894
 TCTTCCAGGTTCACTGAGGGGCTCTGTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTT 586
 CTCACAGCTCTCTGGGACCGGCTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTT 954
 CTCACAGCTCTCTGGGACCGGCTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTT 646
 GACC 962
 GACC 654

31 940 bp mRNA linear EST 16-AUG-2002
 URT 8682031 Lupski_sciatic_nerve Homo sapiens cDNA clone
 6197488 5', mRNA sequence.

31.1 GI:22276239

apiens (human)

apiens
 Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

C http://mgc.nci.nih.gov/.

al Institutes of Health, Mammalian Gene Collection (MGC)

ished (1999)

t: Robert Strausberg, Ph.D.

cgabs-remail.nih.gov

Procurement: Dr. James R. Lupski

Library Preparation: Life Technologies, Inc.

Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

sequencing by: Agencourt Bioscience Corporation

distribution: MGC clone distribution information can be

through the I.M.A.G.E. Consortium/LLNL at:

/image.llnl.gov

LLAM13607 row: j column: 17

uality sequence stop: 453.

Location/Qualifiers

1..940

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:6197488"

/sex="male"

/tissue_type="sciatic nerve"

/dev_stage="adult, 70 Yr"

/lab_host="DH10B"

/clone_lib="Lupski sciatic nerve"

/note="Vector: pCMV-SPORT6 (Life Technologies); Site_1:

Not1; Site_2: SalI; cDNA made by oligo-dT priming.

Directionally cloned using the following adaptors:

5'-TCGACCTCAGGCTCCG-3' and

5'-GACTAGTTCTAGATCGGAGCGGCGCCCT(15)-3'. Size selected >

1 kb for average insert length 1.87 kb. This is a primary

library, non-amplified. Library constructed by Life

Technologies and donated by J. Lupski, M.D./Ph.D. (Baylor

College of Medicine) and is available through Life

Technologies."

31.8%; Score 436; DB 13; Length 940;

arity 100.0%; Pred.No.1e-210;

conservative 0; Mismatches 0; Indels 0; Gaps 0;

301 CAGGACCGCTCGGAATGAAATCCACAGAGAAAGCCAGGATCTCGGCCTTT
 1 CAGGACCGCTCGGAATGAAATCCACAGAGAAAGCCAGGATCTCGGCCTTT
 361 AACGACTAGTTGGCTCGCAGAGTGCACCTTAAGCGCGGAAACACGGGCTCG
 61 AACGACTAGTTGGCTCGCAGAGTGCACCTTAAGCGCGGAAACACGGGCTCG
 421 GCGATCCGACGCCATTAATGAAGTTTCATCCAGCTCGACAGGACGAGGCGGCGG
 121 GCGATCCGACGCCATTAATGAAGTTTCATCCAGCTCGACAGGACGAGGCGGCGG
 481 GTGACCGGACGAGTGCCTGGAGGAAAGCCAGAAATCAACAGCTCCAGCCCTCT
 181 GTGACCGGACGAGTGCCTGGAGGAAAGCCAGAAATCAACAGCTCCAGCCCTCT
 541 TACACCGCCAGATCGGGGAGTTTATAGTACCCGGGCTGGGCTCTACTACCTGTA
 241 TACACCGCCAGATCGGGGAGTTTATAGTACCCGGGCTGGGCTCTACTACCTGTA
 601 CAGGTGCACCTTTGATGAGGGAAGGCTGTCTACCTGAAGCTGGACTTCTGCTGGA
 301 CAGGTGCACCTTTGATGAGGGAAGGCTGTCTACCTGAAGCTGGACTTCTGCTGGA
 661 GTGCTGGCCCTCGCTGCTGGAGGAATTTCTACCCACTCGGCGGCACTTCCCTCGG
 361 GTGCTGGCCCTCGCTGCTGGAGGAATTTCTACCCACTCGGCGGCACTTCCCTCGG
 721 CAGCTCCGCTCTGCC 736
 421 CAGCTCCGCTCTGCC 436

RESULT 17

CF126932

LOCUS

DEFINITION

UI-HF-ETO-avx-k-19-0-UI.r1 NIH_MGC_214 Homo sapiens cDNA c

IMAGE:30563490 5', mRNA sequence.

CF126932

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 697)

Bonaldo, M.F., Lennon, G. and Soares, M.B.

Normalization and subtraction: two approaches to facilitat

discovery

Genome Res. 6 (9), 791-806 (1996)

97044477

8889548

Contact: Soares, MB

Coordinated Laboratory for Computational Genomics

University of Iowa

375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA

Tel: 319 335 8250

Fax: 319 335 9565

Email: bento-soares@uiowa.edu

Tissue Procurement: Mary Hendrix

cDNA Library Preparation: Dr. M. Bento Soares, University

cDNA Library Arrayed by: Dr. M. Bento Soares, University

DNA Sequencing by: Dr. M. Bento Soares, University of Iowa

Clone Distribution: Distribution information can be found

http://genome.uiowa.edu/distribution/humanfl.html

The following repetitive elements were found in this cDN

sequence: 37-143, >GC-rich#Low_complexity

Seq primer: pYX-5.

FEATURES

source

1..697

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:30563490"
/tissue_type="Chondrosarcoma Lung Metastasis cell lines"
/lab_host="DH10B (T1 phage resistant)"
/clone_lib="NIH_MGC_214"
/note="Organ: Lung; Vector: pYX-Asc; Site_1: EcoR I;
Site_2: Not I; The library was constructed according
Bonalido, Lennon and Soares, Genome Research, 6:791-806,
1996. Denatured RNA was size fractionated on a 1% agarose
gel. First strand cDNA synthesis was primed with oligo-dT
primer containing a Not I site. Double strand cDNA was
size selected according to mRNA size fraction, ligated
with EcoR I adaptor, digested with Not I and then cloned
directionally into pYX-Asc vector. The library tag
sequence located between the Not I site and the polyA tail
is GATAGGCCA. Tissue was provided by Mary Hendrix."

31.0%; Score 425; DB 14; Length 697;
rity 100.0%; Pred. No. 4e-205;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
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GCCTGCCAGGAGGAGCTGTGGCAGAGGAGGACCCGTCGGAACCTGAAT 268
GACAGAAAGCCAGGATCTGCGCCCTTTCTGAAACCGACTAGTTCGGCCTGCG 381
GACAGAAAGCCAGGATCTGCGCCCTTTCTGAAACCGACTAGTTCGGCCTGCG 328
TGACCTAAAGCCGGAACACAGGCTCGAAGAGCGATCGACCCATTATGAA 441
TGACCTAAAGCCGGAACACAGGCTCGAAGAGCGATCGACCCATTATGAA 388
TCCACGACCTGGACAGGAGGAGCGAGGAGGTGTGACGGGACAGTGAAGTGC 501
TCCACGACCTGGACAGGAGGAGCGAGGAGGTGTGACGGGACAGTGAAGTGC 448
GGAAGCCAGAAATCAACAGCTCCAGCCCTTCTGCGCTACAAACCGCCAGATCGGGAG 561
GGAAGCCAGAAATCAACAGCTCCAGCCCTTCTGCGCTACAAACCGCCAGATCGGGAG 508
AGTCACCGGGCTGGCTCTACTACCTGTACTGTGTCAGGTGACATTTATGAGGGG 621
AGTCACCGGGCTGGCTCTACTACCTGTACTGTGTCAGGTGACATTTATGAGGGG 568
TGCTACTGAAGCTGACCTGTGCTGGTGGATGTGCTGCGCCCTGCGCTGCGCTG 681
TGCTACTGAAGCTGACCTGTGCTGGTGGATGTGCTGCGCCCTGCGCTGCGCTG 628

686

633

6 834 bp mRNA linear EST 25-SEP-2001
66F1 NIH_MGC_122 Homo sapiens cDNA clone IMAGE:5206217 5',
quence.

6.1 GI:15758344

piens (human)
piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 834)
http://img.ncbi.nih.gov/
l Institutes of Health, Mammalian Gene Collection (MGC)
shed (1999)
: Robert Strausberg, Ph.D.
cgaps-r@mail.nih.gov
Procurement: Life Technologies, Inc.

cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information car
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov

Plate: LLAM11517 row: c column: 18
High quality sequence stop: 772.

FEATURES

Location/Qualifiers
1..834
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5206217"
/lab_host="DH10B"
/clone_lib="NIH_MGC_122"
/note="Organ: pooled lung and spleen; Vector: pCMV
Site 1: NotI; Site 2: EcoRV (destroyed); RNA source
anonymous pool of 24 week female lung, 16 week fem
spleen, and 20-22 week male spleens. Library is c
primed and directionally cloned (EcoRV site is des
upon cloning). Average insert size 1.4 kb. inser
range 1-3 kb. Library is normalized and enriched f
full-length clones and was constructed by C. Gruk
(Invitrogen). Research Genetics tracking code 02
this is a NIH_MGC Library."

ORIGIN

Query Match 30.3%; Score 416; DB 12; Length 834;
Best Local Similarity 99.7%; Pred. No. 1.6e-200;
Matches 656; Conservative 0; Mismatches 0; Indels 2; Gaps
QY 272 CCAGAGGAGCTGTGGCAGAGGAGGACCCGTCGGAACCTGAATCCCCAGT
Db 1 CCAGAGGAGCTGTGGCAGAGGAGGACCCGTCGGAACCTGAATCCCCAGT
QY 332 AAGAAAGCCAGGATCTGCGCCCTTTCTGAAACCGACTAGTTCGGCCTGCGAAGATC
Db 61 AAGAAAGCCAGGATCTGCGCCCTTTCTGAAACCGACTAGTTCGGCCTGCGAAGATC
QY 392 CTAAGGCGGGAACACAGGCTCGAAGAGCGATCGACGCCATTTATGAAGTTCATC
Db 121 CTAAGGCGGGAACACAGGCTCGAAGAGCGATCGACGCCATTTATGAAGTTCATC
QY 452 GACC-TGGACAGGACCGAGCGCAGGCTGTGGACGGGACAGTCAAGTGGCTGGGAC
Db 181 GACCGTGACAGGACCGAGCGCAGGCTGTGGACGGGACAGTCAAGTGGCTGGGAC
QY 511 GCCAGAAATCAACAGCTCCAGCCCTTCTGCGCTACAAACCGCCAGATCGGGAGTTTATP
Db 241 GCCAGAAATCAACAGCTCCAGCCCTTCTGCGCTACAAACCGCCAGATCGGGAGTTTATP
QY 571 ACCCGGGCTGGGCTCTACTACTGTGATGTGTGCTGGCCCTGCGCTGCTGAGGAA
Db 301 ACCCGGGCTGGGCTCTACTACTGTGATGTGTGCTGGCCCTGCGCTGCTGAGGAA
QY 631 TACCTGAAGCTGGACTTCTGTGTGATGTGTGCTGGCCCTGCGCTGCTGAGGAA
Db 361 TACCTGAAGCTGGACTTCTGTGTGATGTGTGCTGGCCCTGCGCTGCTGAGGAA
QY 691 TCAGCCATCGGGCCAGTTCCCTCGGGCCCGCAGCTCCGCTTCGCCAGGTGCTGGC
Db 421 TCAGCCATCGGGCCAGTTCCCTCGGGCCCGCAGCTCCGCTTCGCCAGGTGCTGGC
QY 751 TTGGCCCTCGGGCCAGGCTCTCCCTCGGGATCCGACCCCTCCCTGGGCCCATCTC
Db 481 TTGGCCCTCGGGCCA-GGTCTCTCTCGGGATCCGACCCCTCCCTGGGCCCATCTC
QY 811 GCTGCCCCCTTCTCCTCCTTCTGAGTCTTCCAGGTTCTGAGGGGCCCTTGCTC
Db 540 GCTGCCCCCTTCTCCTCCTTCTGAGTCTTCCAGGTTCTGAGGGGCCCTTGCTC
QY 871 CCACAGTCTGTCAGGCTGCGGGTCCCTCTCCAGAGTCTCTTGGGACACCGGCTCC

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AGTCGTCAGGCTGCGGCTCCCTCGACAGCTCTCTGGGCAACCCGCGTCCCT 657
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96 6.x1 NCI_CGAP_Brn23 Homo sapiens cDNA clone IMAGE:2096962 3',
sequence.
96 413 bp mRNA linear EST 30-MAR-1999
96.1 GI:4268727
apiens (human)
apiens
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
NDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
al Cancer Institute / National Institute of Neurological
ers and Stroke, Brain Tumor Genome Anatomy Project
BTGAP), Tumor Gene Index
ished (1998)
t: Robert Strausberg, Ph.D.
cgapb-re@mail.nih.gov
Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D.,
o, Ph.D.
Library Preparation: M. Bento Soares, Ph.D., M. Fatima
o, Ph.D.
Library Arrayed by: Greg Lennon, Ph.D.
sequencing by: Washington University Genome Sequencing Center
distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
o.llnl.gov/bbrp/image/image.html
Length: 1728 Std Error: 0.00
imer: -40UP from Gibco
quality sequence stop: 410.
Location/Qualifiers
1. .413
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2096962"
/tissue_type="glioblastoma (pooled)"
/lab_host="DH10B"
/clone_lib="NCI CGAP Brn23"
/notes="Organ: Brain; Vector: pTT73D-Pac (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTACCATCTGAAGTGGGAGCGCGCCATATCTTTTCTTTTCTTTTCTTTT
T 3']; double-stranded cDNA was ligated to Eco RI
adaptors (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of the modified pTT73 vector.
Library is normalized, and was constructed by Bento
Soares and M.Fatima Bonaldo."
28.7%; Score 394; DB 9; Length 413;
arity 100.0%; Pred. No. 2.4e-189;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
|CTGGGCGCTGTCAGCTGTTTCCATCCACATATACATATCCCACTCTTAT 1039
|CCTGGGCGCTGTTTCCAGTGTTCATCCACATATATACATATCCCACTCTTAT 354
|CAACTCCCCACCGCCCACTCTCCACCTCCTAGCTCCCACTCCCACTCCCTTTG 1099
|||||
|CAACTCCCCACCGCCCACTCTCCACCTCCTAGCTCCCACTCCCACTCCCTTTG 294
|||||
|CCCCAGTATCTGACTCCCCCTGGGCCACAGACCCCGAGGCGATGTGTTCACGT 1159
|CCCCAGTATCTGACTCCCCCTGGGCCACAGACCCCGAGGCGATGTGTTCACGT 234
|CTGTGGGAGGATGGTCCAGAACACCCCACTTCAGGCACCTAAGAGGGGCTGGAC 1219
```

```
Db 233 TACTCTGTGGGCAAGGATGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGCT
|||||
Oy 1220 CTGGCGGCAAGGACCAAGAGAGACTGGGCGCTAGGCCAGGAGTTCCCAAAATGTGAGG
|||||
Db 173 CTGGCGGCAAGGACCAAGAGAGACTGGGCGCTAGGCCAGGAGTTCCCAAAATGTGAGG
|||||
Oy 1280 AGAAACAAGACAGAGCTCTCTCCCTTGAGAAATCCCTGTGGATTTTAAACACAGATAT
|||||
Db 113 AGAAACAAGACAGAGCTCTCTCCCTTGAGAAATCCCTGTGGATTTTAAACACAGATAT
|||||
Oy 1340 TTATTATTATTGTGACAAATGTTGATAATGG 1373
|||||
Db 53 TTATTATTATTGTGACAAATGTTGATAATGG 20
|||||
RESULT 20
BM971606/c
LOCUS
DEFINITION
UI-CF-ECI-abl-p-06-0-UI.s1 UI-CF-ECI Homo sapiens cDNA clc
UI-CF-ECI-abl-p-06-0-UI 3', mRNA sequence.
ACCESSION
BM971606
VERSION
BM971606.1 GI:19589193
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 568)
Bonaldo,M.F., Lennon,G. and Soares,M.B.
Normalization and subtraction: two approaches to facilitat
discovery
Genome Res. 6 (9), 791-806 (1996)
JOURNAL
MEDLINE
PUBMED
8889548
COMMENT
Contact: McCray, PB
McCray Lab
University of Iowa
2024 University of Iowa Med Labs, Iowa City, IA 52242, USA
Tel: 319 356 4866
Fax: 319 356 7171
Email: paul-mccray@uiowa.edu
Tissue Procurement: Dr. M. J. Welsh, University of Iowa
cDNA Library Preparation: Dr. M. Bento Soares, University
cDNA Library Arrayed by: Dr. M. Bento Soares, University
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Researchers may obtain clones from Re
Genetics (www.resgen.com) or from Open Biosystems
(www.openbiosystems.com).
The following repetitive elements were found in this cDN
sequence: 1-82, >AT-rich#low_complexity (matched complim
Seq primer: M13 FORWARD
POLYA=Yes.
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FEATURES
Location/Qualifiers
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/clone_lib="UI-CF-ECI"
/notes="Organ: Lung; Vector: pTT73-Pac (Pharmacia)
modified polylinker; Site 1: Eco RI; Site 2: Not
UI-CF-ECI is a normalized cDNA library containing
following tissue(s): Normal lung from adult and
day 64, day 87, week 19 and week 42. The library
constructed according to Bonaldo, Lennon and Soa
Genome Research, 6:791-806, 1996. First strand cl
synthesis was primed with an oligo-dT primer cont
Not I site. Double stranded cDNA was ligated to
adaptor, digested with Not I, and cloned directic
```

into pT73-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (GT)₁₈ tail. The sequence tag for this library is AAGTGGCTTAC.
TAG TISSUE=Normal Lung Epithelial Cells Tissue nos 369-371 and 380-383
TAG LIB=UI-CF-EC1
TAG_SEQ=AAGTGGCTTAC"

28.7%; Score 394; DB 12; Length 568;
rity 100.0%; Pred. No. 2.5e-189;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
CTGGGCGTTCACGTGTTTCCATCCACATAAATACAGTATCCCACTTTAT 1039
CTGGGCGTTCACGTGTTTCCATCCACATAAATACAGTATCCCACTTTAT 362
AACTCCCCCAGCGCCACTCTCCACCTCAGTCTCCCAATCCCTGACCCCTTG 1099
AACTCCCCCAGCGCCACTCTCCACCTCAGTCTCCCAATCCCTGACCCCTTG 302
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GGCAGGAGCCAAAGAGACTGGGCTAGGCCAGGAGTTCCCAATGTGAGGGGCG 122
CAAGACAAGCTCCCTCCCTTGAGAAATTCCTGTGGAATTTTAAACAGATATT 1339
CAAGACAAGCTCCCTCCCTTGAGAAATTCCTGTGGAATTTTAAACAGATATT 62
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TATTATTGTGACAAATGTTGATAAATGG 28

4 569 bp mRNA linear EST 23-SEP-2002
1-bdu-c-24-0-UI.s1 NCI CGAP FE1 Homo sapiens cDNA clone
1-bdu-c-24-0-UI 3', mRNA sequence.
4
4.1 GI:23298519
piens (human)
piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 569)
P http://www.ncbi.nlm.nih.gov/ncicgap.
1 Cancer Institute, Cancer Genome Anatomy Project (CGAP),
ene Index
shed (1997)
: Robert Strausberg, Ph.D.
cgapbs-remail.nih.gov
Procurement: James Martin
Library preparation: Dr. M. Bento Soares, University of Iowa
Library Arrayed by: Dr. M. Bento Soares, University of Iowa
Sequencing by: Dr. M. Bento Soares, University of Iowa
Distribution: Clone distribution information can be obtained
from: M. Bento Soares, bento-soares@uiowa.edu
allowing repetitive elements were found in this cDNA.
re: 1-82, >AT-rich/low complexity (matched complement)
mer: M13 FORWARD
res.

Location/Qualifiers
1. 569
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="UI-H-FE1-bdu-c-24-0-UI"
/tissue_type="Cell lines"
/dev_stage="Adult"
/lab_host="DH10B (Life Technologies)"
/clone_lib="NCI CGAP FE1"
/note="Organ: Chondrosarcoma; Vector: pT73-Pac (Pharmacia) with a modified polylinker; Site 1: Ec Site 2: Not I; NCI CGAP FE1 is a normalized cDNA 1 derived from a pool of mRNA obtained from 3 cell 1 from grade II chondrosarcoma tissues. The library constructed according to Bonaldo, Lennon and Soares Genome Research, 8:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer conta Not I site. Double stranded cDNA was ligated to a adaptor, digested with Not I, and cloned director into pT73-Pac vector. The oligonucleotide used to the synthesis of first-strand cDNA contains a libi sequence that is located between the Not I site an (GT)₁₈ tail. The sequence tag for this library is CCGTACGGAC. The cell lines were provided by Dr Jan Martin from the University of Iowa.
TAG TISSUE=Human grade 2 chondrosarcoma cell line
TAG LIB=UI-H-FE1
TAG_SEQ=CGCTACGGAC"

ORIGIN

Query Match . 28.6%; Score 393; DB 13; Length 569;
Best Local Similarity 99.8%; Pred. No. 8.2e-189;
Matches 443; Conservative 0; Mismatches 1; Indels 0; Gaps
QY 930 TGCCCAACCTCAGCGCTTTTGTCTCCAGACCTGCCCCCTCCCTCTAGAGGCTGCC
DB 471 TGCCCAACCTCAGCGCTTTTGTCTCCAGACCTGCCCCCTCCCTCTAGAGGCTGCC
QY 990 CCGTGTTCAGTGTGTTTCCATCCACATAAATACAGTATCCCACTTTATCTTACAT
DB 411 CCGTGTTCAGTGTGTTTCCATCCACATAAATACAGTATCCCACTTTATCTTACAT
QY 1050 CCCCACCGCCCACTCTCCACCTCAGTCTCCCCCAATCCCTGACCCCTTTGAGGCGCC
DB 351 CCCCACCGCCCACTCTCCACCTCAGTCTCCCCCAATCCCTGACCCCTTTGAGGCGCC
QY 1110 TGAATCTGACTCCCCCTGGCCACAGACCCCGAGGCATTGTTCACGTACTCT
DB 291 TGAATCTGACTCCCCCTGGCCACAGACCCCGAGGTCATTGTGTTCACTGTACTCT
QY 1170 GCAAGGATGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGCTGGACCTGGCG
DB 231 GCAAGGATGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGCTGGACCTGGCG
QY 1230 GAAGCCAAAGAGACTGGGCTAGGCCAGGAGTTCCCAATGTGAGGGGCGAGAAAC
DB 171 GAAGCCAAAGAGACTGGGCTAGGCCAGGAGTTCCCAATGTGAGGGGCGAGAAAC
QY 1290 CAACTCTCCCTTCAGAAATTCCTGTGGAATTTTAAACAGATATTATTTTATT
DB 111 CAACTCTCCCTTCAGAAATTCCTGTGGAATTTTAAACAGATATTATTTTATT
QY 1350 TTGTGACAAATGTTGATAAATGG 1373
DB 51 TTGTGACAAATGTTGATAAATGG 28

RESULT 22
BI824443
LOCUS
DEFINITION
60308693F1 NIH_MGC_115 Homo sapiens cDNA clone IMAGE:5179:
mRNA sequence.

13
13.1 GI:15935993
apiens (human)
apiens
ata; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
ses 1 to 531)
2 http://mgc.nci.nih.gov/
al Institutes of Health, Mammalian Gene Collection (MGC)
ished (1999)
t: Robert Strausberg, Ph.D.
cgapbs-remail.nih.gov
Library Preparation: Life Technologies, Inc.
Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
sequencing by: Inocyte Genomics, Inc.
distribution: MGC clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
/image.llnl.gov
LLM11447 row: j column: 23
uality sequence stop: 529.
Location/Qualifiers
1. .531
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5179510"
/lab_host="DH10B"
/clone_lib="NIH_MGC_115"
/note="Organ: pooled brain, lung, testis; Vector:
pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA
source anonymous pool of 6 male brains, age range 23-27; 1
male lung, age 27; and 1 male testis, age 69. Library is
oligo-dT primed and directionally cloned (EcoRV site is
destroyed upon cloning). Average insert size 1.8 kb,
insert size range 1-3 kb. Library is normalized and
enriched for full-length clones and was constructed by C.
Gruber (Invitrogen). Research Genetics tracking code
021. Note: this is a NIH_MGC Library."
27.3%; Score 375; DB 12; Length 531;
arity 99.8%; Pred. No. 1.2e-179;
onservative 0; Mismatches 1; Indels 0; Gaps 0;
CTCCCTCGGATCGCAGCTCTCCCTCGGCGCCATCTCAGCTGCCCTTCCTCA 826
CTCCCTCGGATCGCAGCTCTCCCTCGGCGCCATCTCAGCTGCCCTTCCTCA 165
CTTCGGACTCTTCAGGTTCACTGAGGGCCCTGCTCCCGCAGTCTCCAGG 886
CTTCGGACTCTTCAGGTTCACTGAGGGCCCTGCTCCCGCAGTCTCCAGG 225
CGGCTCCCTCGCAGCTCTCTGCGGACCCCGTCCCTCTGCCACCTCAGCGG 946
CGGCTCCCTCGCAGCTCTCTGCGGACCCCGTCCCTCTGCCACCTCAGCGG 285
TTGCTCAGAGCTGCCCTCCCTCTAGAGGCTGCTGGGCTGTTCAGTGTTC 1006
TTGCTCAGAGCTGCCCTCCCTCTAGAGGCTGCTGGGCTGTTCAGTGTTC 345
XCACATAATACAGTATTCACACTCTTATTTACAACTCCCGCCAGCTCTC 1066
XCACATAATACAGTATTCACACTCTTATTTACAACTCCCGCCAGCTCTC 405
XCACATAATACAGTATTCACACTCTTATTTACAACTCCCGCCAGCTCTC 1126
XCACATAATACAGTATTCACACTCTTATTTACAACTCCCGCCAGCTCTC 465
XCACAGACCCAGGCGATGTTCACGTACTGTGGGCAAGATGGGTCCAGA 1186
XCACAGACCCAGGCGATGTTCACGTACTGTGGGCAAGATGGGTCCAGA 525

QY 1187 AGACCC 1192
Db 526 AGACCC 531
RESULT 23
CB998034
LOCUS
DEFINITION
CB998034 824 bp mRNA linear EST 01
IMAGE:30348032 5', mRNA sequence.
CB998034
CB998034.1 GI:30292554
EST.
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 824)
NIH-MGC http://mgc.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Dr. Stefan Hansson
cDNA Library Preparation: Michael J. Brownstein (NHGRI) w
and advice from Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information ca
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: NDAM390 row: b column: 09
High quality sequence stop: 415.
Location/Qualifiers
1. .824
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:30348032"
/tissue_type="pre-eclamptic placenta"
/lab_host="DH10B Tona"
/clone_lib="NIH_MGC_148"
/note="Organ: placenta; Vector: pBluescriptR; Sit
all-XhoI; Site 2: BamH; Library is oligo-dT prime
directionally cloned using primer
5'-TTTTTTTTTTTTTTVN-3', size selected for avera
size 2.3 kb and normalized to ROT 5. This is a f
library enriched for full-length clones and cons
using the Cap-trapper method (Carninci, in prepa
Library constructed by M. Brownstein (NIH/NHGRI
National Institutes of Health). Note: this is a N
Library."
Query Match 26.4%; Score 362; DB 14; Length 824;
Best Local Similarity 99.8%; Pred. No. 5.9e-173;
Matches 412; Conservative 0; Mismatches 1; Indels 0; G
QY 52 TCCCTCGGCTCCCGGATGGGGGGCGGTGAGGCGAGCAGCCCGCCGCCAI
Db 70 TCCCTCGGCTCCCGGATGGGGGGCGGTGAGGCGAGCAGCCCGCCGCCAI
QY 112 GCCCGTTCGAGCAGAGCGGAGGGGGCGCGGGGGGAGCGGGCAGCCGCTCT
Db 130 GCCCGTTCGAGCAGAGCGGAGGGGGCGCGGGGGGAGCGGGCAGCCGCTCT
QY 172 CCGCTCGGCTTCGGCTTCGGCTTCGGCTTCGGCTTCGGCTTCGGCTTCGGCT
Db 190 CCGCTCGGCTTCGGCTTCGGCTTCGGCTTCGGCTTCGGCTTCGGCTTCGGCT
QY 232 AGTTTGGGAGCCCGGGATCGCTCTCCGCCAGGAGCTGCCCGAGGAGCTGGT

3GGAGCGCGGCGATCGGTCTCGGCCAGGAGCGCTGCCAGGAGCGTGGTGGCA 309
 3GACGAGCGCGGCGAACTGAATCCCGACAGAGAGAGAGAGAGAGAGAGAGAG 351
 3GACGAGCGCGGCGGAACTGAATCCCGACAGAGAGAGAGAGAGAGAGAGAGAG 369
 CCTGAACCGAGCTAGTTCGGGCTCGCAGAAGTGACCTTAAAGGCGGGAACACGG 411
 CCTGAACCGAGCTAGTTCGGGCTCGCAGAAGTGACCTTAAAGGCGGGAACACGG 429
 AAGAGCGATCGACCGCATTTATGAGTTTCATCCAGACCTGGACAGGA 464
 AAGAGCGATCGACCGCATTTATGAGTTTCATCCAGACCTGGACAGGA 482

9 440 bp mRNA linear EST 12-MAR-2002
 .Y1 Melton Normalized Human Islet 4 N4-HIS 1 Homo sapiens
 one IMAGE:5675843 5' similar to TR:054907 054907 TNF-RELATED
 DUCER OF APOPTOSIS 1, mRNA sequence.

9.1 GI:17122611

piens (human)

piens
 ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 a; Eutheria; Primates; Catarrhini; Hominidae; Homo.

es 1 to 440)
 D., Brown, J., Kenty, G., Permutt, A., Lee, C., Kaestner, K.,
 a, I., Scearce, M., Brestelli, J., Gradwohl, G., Clifton, S.,
 L., Marra, M., Pape, D., Wylie, T., Martin, J., Blistain, A.,
 A., Theising, B., Ritter, E., Ronko, I., Bennett, J.,
 S.M., Gibbons, M., McCann, R., Cole, R., Tsagaris, V., R.,
 S.T., Jackson, Y., and Bowers, Y.
 ne Pancreas Consortium
 shed (2000)

: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
 ne Pancreas Consortium

University, Howard Hughes Medical Institute
 Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
 8

7-495-1812
 7-495-8557
 dmeltont@bioph.harvard.edu

was constructed by Dr. Douglas Melton DNA sequencing by:
 ng a clone please contact: Juliana Brown
 fas.harvard.edu) This sequence now available from the IMAGE
 ium, for clone orders contact: info@image.llnl.gov
 ality sequence stop: 415.
 Location/Qualifiers

1. .440
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:5675843"
 /sex="Both"
 /tissue_type="Islets of Langerhans"
 /dev_stage="Adult"
 /lab_host="DH10B"

/notes="Organ: Pancreas; Vector: pSPORT1; Site 1: Not 1;
 Site 2: Sal 1; Starting library constructed using
 SuperScript Plasmid Library kit (Life Technologies). cDNA
 made by oligo-dT priming. Size-selected by column
 fractionation; average insert size 1.08 kb. Library was
 amplified once on solid support and plasmid DNA from
 library was prepared. The library DNA was normalized by
 method #4 from Bionano, Lennon, and Soares 1996 Genome
 Research 6:791-806; 0.5 microgram single-stranded library
 plasmid DNA was mixed with 5 micrograms PCR product
 representing library inserts and hybridized to an EcoT

20. Single-stranded (unhybridized) plasmids were
 by hydroxyapatite chromatography and used to make
 library."

Query Match 25.6%; Score 352; DB 12; Length 440;
 Best Local Similarity 99.8%; Pred. NO. 6.6e-168;
 Matches 402; Conservative 0; Mismatches 1; Indels 0; Gs

ORIGIN

QY 808 AAGGTGCCCCCTTCCTCAGCTACTTCCAGGTTCTCCAGGTTCTACTGAGGGCCCTG
 DB 38 AAGGTGCCCCCTTCCTCAGCTACTTCCAGGTTCTACTGAGGGCCCTG
 QY 868 TCCCCACAGTGTCTCCAGGCTGCCGGCTCCCTCGACAGCTCTCTGGGACCCCGGTC
 DB 98 TCCCCACAGTGTCTCCAGGCTGCCGGCTCCCTCGACAGCTCTCTGGGACCCCGGTC
 QY 928 TGTGCCCCACCTCAGCGCTCTTTGCTCCAGACCTGCCCTCCCTCTAGAGGCTGC
 DB 158 TGTGCCCCACCTCAGCGCTCTTTGCTCCAGACCTGCCCTCCCTCTAGAGGCTGC
 QY 988 GGCCTGTTTCAGTGTTTCCATCCACATAATACAGTATCCCACTCTTATCTTAC
 DB 218 GGCCTGTTTCAGTGTTTCCATCCACATAATACAGTATCCCACTCTTATCTTAC
 QY 1048 TCCCCCACCGCCCACTCTCCACCTCACTAGTCTCCCAATCCCTGACCCCTTTGAGGCG
 DB 278 TCCCCCACCGCCCACTCTCCACCTCACTAGTCTCCCAATCCCTGACCCCTTTGAGGCG
 QY 1108 AGTGATCTCGACTCCCTCCCTGSCCAGACCCCGGAGGATGTTGTTACTTACTC
 DB 338 AGTGATCTCGACTCCCTCCCTGSCCAGACCCCGGAGGATGTTGTTACTTACTC
 QY 1168 GGGCAAGGATGGTCCAGAGACCCCACTCAGGCACTAAGAG 1210
 DB 398 GGGCAAGGATGGTCCAGAGACCCCACTCAGGCACTAAGAG 440

RESULT 25

B0674188

LOCUS

DEFINITION

BO674188

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BO674188

AGENCY

BO674188

BO674188

BO674188

BO674188

BO674188

BO674188

BO674188

BO674188

BO674188

BO674188

BO674188

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BO674188

BO674188

/clone_lib="NIH_MGC_102"
 /note="Organ: salivary gland; Vector: pOTB7; Site_1: XhoI;
 Site_2: EcoRI; cDNA made by oligo-dT priming.
 Directionally cloned into EcoRI/XhoI sites using the
 following 5' adaptor: GGCACGAG(G). Library constructed
 by Ling Hong in the laboratory of Gerald M. Rubin
 (University of California, Berkeley) using ZAP-cDNA
 synthesis kit (Stratagene) and Superscript II RT (Life
 Technologies). Note: this is a NIH_MGC Library."

25.4%; Score 349; DB 13; Length 951;
 arity 100.0%; Pred. No. 2.6e-166;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CGGCTCGCAGAGTGCACCTAAAGCGCGAAACACGGCTCGAAGCGATCGC 428
 CGGCTCGCAGAGTGCACCTAAAGCGCGAAACACGGCTCGAAGCGATCGC 60
 CATTATGAAGTTTCATCCACGACCTGGACAGGACGCGCAGGCGAGTGTGACGG 488
 CATTATGAAGTTTCATCCACGACCTGGACAGGACGCGCAGGCGAGTGTGACGG 120
 GTGAGTGGCTGGAGGAGGACAGATCAACAGCTCCAGCCCTCTCGCTACAAACGG 548
 GTGAGTGGCTGGAGGAGGACAGATCAACAGCTCCAGCCCTCTCGCTACAAACGG 180
 ATCCGGGAGTTTATAGTCAACCGGGCTGGCTCTACTACCTGTACTGTGAGTGCA 608
 ATCCGGGAGTTTATAGTCAACCGGGCTGGCTCTACTACCTGTACTGTGAGTGCA 240
 GATGAGGGGAGGCTGTCTACTGAAGCTGGACTTGTGTGTGATGTGTCTGCGC 668
 GATGAGGGGAGGCTGTCTACTGAAGCTGGACTTGTGTGTGATGTGTCTGCGC 300
 CGCTGCTCGGAGGATTTCTAGCCACTCGGCCAGTTCCCTCGGG 717
 CGCTGCTCGGAGGATTTCTAGCCACTCGGCCAGTTCCCTCGGG 349

16 377 bp mRNA linear EST 15-FEB-2002
 5.Y1 Human insulinoma Homo sapiens cDNA 5', mRNA sequence.
 16
 16.1 GI:18680159

sapiens (human)
 Meta; Chordata; Craniata; Vertebrata; Euteleostomi;
 ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 to 377)
 D., Brown, J., Kenty, G., Permut, A., Lee, C., Kaestner, K.,
 ka, I., Scarce, M., Brestelli, J., Gradwohl, G., Clifton, S.,
 T., L., Marra, M., Pape, D., Wylie, T., Martin, J., Bliscain, A.,
 T., A., Theising, B., Ritter, E., Ronko, I., Bennett, J.,
 as, M., Gibbons, M., McCann, R., Cole, R., Teagareishvili, R.,
 ms, T., Jackson, Y., and Bowers, Y.
 ine Pancreas Consortium
 ished (2000)
 ESTs: ih15605.x1
 it: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
 ine Pancreas Consortium
 d University, Howard Hughes Medical Institute
 of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
 138
 317-495-1812
 317-495-8557
 : dmelton@biohp.harvard.edu
 cy was constructed by Dr. J. J. Ferrer in vivo mass-excised to
 script SK- by Dr. H. Inoue DNA sequencing by: Washington
 sity Genome Sequencing Center for information on obtaining a
 please contact: Dr. Hiroshi Inoue (hinoue@im.wustl.edu)

Seq primer: -40RP from Gibco.
 Location/Qualifiers
 1. 377
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /tissue_type="insulinoma"
 /lab_host="DH10B (phage-resistant)"
 /clone_lib="Human insulinoma"
 /note="Organ: pancreas; Vector: pBluescript SK-;
 XhoI; Site_2: EcoRI; Constructed with lambda ZAPI
 (Stratagene) by Dr. J. J. Ferrer, in vivo mass-excise
 pBluescript SK- by Dr. H. Inoue following the Wa
 University protocol
 (http://genome.wustl.edu/est/lambda_protocol.shtm
 please contact Hiroshi Inoue, MD/PhD for further
 information on this library (Metabolism Division
 Laboratory, Washington University School of Medi
 8127, 660 S Euclid Ave, St. Louis, MO 63110). NC
 is a Washington University Pancreas EST project 1

ORIGIN

Query Match 24.8%; Score 341; DB 12; Length 377;
 Best Local Similarity 100.0%; Pred. No. 2.6e-162; Indels 0; C
 Matches 341; Conservative 0; Mismatches 0; Indels 0; C
 QY 1033 CTCCTATCTTACAACTCCCGCCACCGCCACTCTCCACCTCACTAGTCCCAATCC
 DB 8 CTCCTATCTTACAACTCCCGCCACCGCCACTCTCCACCTCACTAGTCCCAATCC
 QY 1093 CCCTTTGAGGCCCCAGTGATCTCGACTCCCGCCCTGGCCACAGACCCCGAGGCA
 DB 68 CCCTTTGAGGCCCCAGTGATCTCGACTCCCGCCCTGGCCACAGACCCCGAGGCA
 QY 1153 TTCACTGTACTCTGTGGCAAGGATGGTCCAGAAAGACCCCACTTCAGGCACTAAC
 DB 128 TTCACTGTACTCTGTGGCAAGGATGGTCCAGAAAGACCCCACTTCAGGCACTAAC
 QY 1213 GCTGGACCTGGCGCAGGAGCAAGCAAGAGACTGGGCTTAGCCAGGAGTCCCAAF
 DB 188 GCTGGACCTGGCGCAGGAGCAAGCAAGAGACTGGGCTTAGCCAGGAGTCCCAAF
 QY 1273 AGGGCGGAGAAACAGACAAAGCTCCCTCCCTGAGAAATTCCTGTGATTTTAAAT
 DB 248 AGGGCGGAGAAACAGACAAAGCTCCCTCCCTGAGAAATTCCTGTGATTTTAAAT
 QY 1333 TATTATTTTATTATTATTGTGACAAATGTTGATAAATGG 1373
 DB 308 TATTATTTTATTATTATTGTGACAAATGTTGATAAATGG 348

RESULT 27

BE858778 710 bp mRNA linear EST 2;
 BE858778/c LOCUS
 7f9b06.x1 NCI_CGAP_Brn23 Homo sapiens cDNA clone IMAGE:3;
 DEFINITION
 similar to contains element MER32 repetitive element ; mi
 sequence.
 ACCESSION
 BE858778
 VERSION
 BE858778.1 GI:10374165
 KEYWORDS
 EST.
 SOURCE
 Homo sapiens (human)
 ORGANISM
 Homo sapiens
 Eukaryota; Chordata; Craniata; Vertebrata; Eutele
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 710)
 REFERENCE
 NCI/NINDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute / National Institute of Neurolo
 Disorders and Stroke, Brain Tumor Genome Anatomy Project
 (CGAP/BTGAP), Tumor Gene Index
 Unpublished (1998)
 JOURNAL
 COMMENT
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenf

Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Ph.D.
Library Arrayed by: Greg Lennon, Ph.D.
Sequencing by: Washington University Genome Sequencing Center
Distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL, send email to:
age.llnl.gov
mer: -40UP from Gibco
ality sequence stop: 342.
Location/Qualifiers
1. .710

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3304691"
/tissue_type="glioblastoma (pooled)"
/lab_host="DH10B"
/clone_lib="NCI CGAP Brn23"
/note="Organ: Brain; Vector: pTVT3D-Pac (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo (dT) primer [5'
TGTACCAATCTGAAGTGGAGCGCGCATATCTTTTTTTTTTTTTTTTTT
T 3']; double-stranded cDNA was ligated to Eco RI
adaptors (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of the modified pTVT3 vector.
Library is normalized, and was constructed by Bento
Soares and M.Fatima Bonaldo."

24.0%; Score 329; DB 10; Length 710;
rity 100.0%; Pred. No. 46-156;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
CCCCACGCCCACTCTCCACTACTAGCTCCCAATCCCTGACCTTTGAGGCC 1104
CCCCACGCCCACTCTCCACTACTAGCTCCCAATCCCTGACCTTTGAGGCC 289
TGATCTGACTCCCCCTGGCCACAGACCCCGAGGCGATTGTCTACTGTACTC 1164
TGATCTGACTCCCCCTGGCCACAGACCCCGAGGCGATTGTCTACTGTACTC 229
ACAGGATGGTCCAGAGACCCCACTTACGACCTAAGAGGGGCTGACCTGGC 1224
ACAGGATGGTCCAGAGACCCCACTTACGACCTAAGAGGGGCTGACCTGGC 169
CAAGCCAAAGAGAGCTGGCCCTAGGCGAGGATCCCAATGTGAGGGCGAGAAA 1284
CAAGCCAAAGAGAGCTGGCCCTAGGCGAGGATCCCAATGTGAGGGCGAGAAA 109
CAAGCTCTCTCCCTTGAGAAATCCCTGTGGATTCTTTAAACAGATATTTTAT 1344
CAAGCTCTCTCCCTTGAGAAATCCCTGTGGATTCTTTAAACAGATATTTTAT 49
ATTGTACAAATATTGTGATAATGG 1373
ATTGTACAAATATTGTGATAATGG 20

367 bp mRNA linear EST 27-FEB-2002
1l-acs-a-05-0-UI.81 UI-E-CQ1 Homo sapiens cDNA clone
1l-acs-a-05-0-UI 3', mRNA sequence.
2.1 GI:18967291
apiens (human)
apiens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ses 1 to 367)
M.F., Lennon, G. and Soares, M.B.

TITLE Normalization and subtraction: two approaches to facilitate
discovery
JOURNAL Genome Res. 6 (9), 791-806 (1996)
MEDLINE 97044477
PubMed 8889548
COMMENT Contact: Soares, MB
Coordinated Laboratory for Computational Genomics
University of Iowa
375 Newton Road 4156 MEBRF, Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: bento-soares@uiowa.edu
Tissue Procurement: Dr. Gregg Hageman
cDNA Library Preparation: Dr. M. Bento Soares, University of
Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Researchers may obtain clones from Res
Genetics (www.resgen.com).
The following repetitive elements were found in this cDNA
sequence: 1-94, >At rich#Low complexity (matched complement
Seq primer: M13 Forward
POLYA=Yes.

FEATURES
source

Location/Qualifiers
1. .367
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="UI-E-CQ1-acs-a-05-0-UI"
/tissue_type="optic nerve"
/dev_stage="adult"
/clone_lib="UI-E-CQ1"
/note="Organ: eye; Vector: pTVT3-Pac (Pharmacia) v
modified polylinker; Site 1: EcoRI; Site 2: Not I
UI-E-CQ1 is a normalized cDNA library containing t
following tissue(s): optic nerve. The library was
constructed according to Bonaldo, Lennon and Soar
Genome Research, 6:791-806, 1996. First strand cD
Synthesis was primed with an oligo-dT primer con
Not I site. Double stranded cDNA was ligated to a
adaptor, digested with Not I, and cloned directio
into pTVT3-Pac vector. The oligonucleotide used t
the synthesis of first-strand cDNA contains a lib
sequence that is located between the Not I site a
(dT)18 tail. The sequence tag for this library is
CCATTAAAGTG. This library was created for the prog
discovery in the Visual System, supported by Natio
Institute (NEI).
TAG_TISSUE=human optic nerve
TAG_LIB=UI-E-CQ1
TAG_SEQ=CCATTAAAGTG"

ORIGIN

Query Match 23.9%; Score 328; DB 12; Length 367;
Best Local Similarity 100.0%; Pred. No. 11e-155;
Matches 328; Conservative 0; Mismatches 0; Indels 0; G
QY 1045 ACTCCCCACCGCCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCTTTGAG
DB 367 ACTCCCCACCGCCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCTTTGAG
QY 1106 CCAGTGATCTGACATCCCTCCCTGGCCACAGACCCCGAGGCGATTGTCTCACTGTAT
DB 307 CCAGTGATCTGACATCCCTCCCTGGCCACAGACCCCGAGGCGATTGTCTCACTGTAT
QY 1166 GTGGGCAAGATGGGTCCAGAGAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTT
DB 247 GTGGGCAAGATGGGTCCAGAGAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTT
QY 1226 GCAGGAGCAAGAGAGACTGGGCGCTAGGCGAGGAGTCCCAATGTGAGGGCGAG
DB 187 GCAGGAGCAAGAGAGACTGGGCGCTAGGCGAGGAGTCCCAATGTGAGGGCGAG


```
CAAGCTCCTCCCTTGAGAAATCCCTGCGAATTTTAAACAGATATATTTTATT 1345
|||||
CAAGCTCCTCCCTTGAGAAATCCCTGCGAATTTTAAACAGATATATTTTATT 68
|||||
ITGTGACAAATGTTGATAAATGG 1373
|||||
ITGTGACAAATGTTGATAAATGG 40
|||||

22      346 bp      mRNA      linear      EST 29-SEP-2000
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sequence.
22
22.1 GI:10374253
apiens (human)
apiens
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ses 1 to 346)
NDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
al Cancer Institute / National Institute Of Neurological
ars and Stroke, Brain Tumor Genome Anatomy Project
BTGP), Tumor Gene Index
ished (1998)
t: Robert Strausberg, Ph.D.
cgapbs-r@mail.nih.gov
: Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D.,
Library Preparation: M. Bento Soares, Ph.D., M. Fatima
o, Ph.D.
Library Arrayed by: Greg Lennon, Ph.D.
sequencing by: Washington University Genome Sequencing Center
distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL, send email to:
mage.llnl.gov
amer: -40UP from Gibco
quality sequence stop: 344.
Location/Qualifiers
1. 346
/organism="Homo sapiens"
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/tissue_type="glioblastoma (pooled)"
/lab_host="DH10B"
/clone_lib="NCI_CGAP_Brn23"
/notes="Organ: brain; Vector: pTT73D-Pac (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5',
TGTTACCAATCTGAAGTGGAGCGCGCATATCTTTTATTTTATTTTATTTT
T 3']; double-stranded cDNA was ligated to Eco RI
adaptors (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of the modified pT73 vector.
Library is normalized, and was constructed by Bento
Soares and M.Fatima Bonaldo."
```

23.8%; Score 327; DB 10; Length 346;

arity 100.0%; Pred.No. 3.6e-155; Mismatches 0; Indels 0; Gaps 0;

CCCCACGGCCACTCTCCACTAGTCTCCCAATCCCTGACCCCTTTGAGGCCCC 1106

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CCCCACGGCCACTCTCCACTAGTCTCCCAATCCCTGACCCCTTTGAGGCCCC 287

|||||

TGATCTGACTCCCTCGCCACAGACCCCGAGGCAATGTGTTCACTGTACTCTG 1166

|||||

TGATCTGACTCCCTCGCCACAGACCCCGAGGCAATGTGTTCACTGTACTCTG 227

|||||

3CAGGATGGGTCCAGNAGACCCCACTTCAGGCACTAAGAGGGGCTGGACCTGGCG 1226

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Db      226      TGGCAAGGATGGTCCGAAGACCCCACTTCAGGCACTAAGAGGGCTGGACCTG
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Qy      1227      CAGGAAGCCAAAGAGACTGGGCGCTAGGCCAGGAGTTCCTCCAAATGTGAGGGCGGAGA
|||||
Db      166      CAGGAAGCCAAAGAGACTGGGCGCTAGGCCAGGAGTTCCTCCAAATGTGAGGGCGGAGA
|||||
Qy      1287      AGCAAGCTCTCTCCCTTGAGAAATCCCTGCGAATTTTAAACAGATATATTTTATT
|||||
Db      106      AGCAAGCTCTCTCCCTTGAGAAATCCCTGCGAATTTTAAACAGATATATTTTATT
|||||
Qy      1347      TTATTGTGACAAATGTTGATAAATGG 1373
|||||
Db      46      TTATTGTGACAAATGTTGATAAATGG 20
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RESULT 30
AW195034/c
LOCUS
DEFINITION
x45912.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:26
mRNA sequence.

ACCESSION
AW195034
VERSION
AW195034.1 GI:6474026
KEYWORDS
EST.
SOURCE
Homo sapiens (human)

ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 337)

REFERENCE
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS
National Cancer Institute, Cancer Genome Anatomy Project (

TITLE
Tumor Gene Index
JOURNAL
Unpublished (1997)

COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Mic
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Sequencing by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencin
Clone distribution: NCI-CGAP clone distribution informati
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco
High quality sequence stop: 330.

Location/Qualifiers
1. 337

FEATURES
source
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/clone="IMAGE:2696710"
/lab_host="DH10B"

/clone_lib="NCI_CGAP_Kid11"
/note="Organ: kidney; Vector: pT73D-Pac (Pharma
a modified polylinker; Site 1: Not I; Site 2: Eco
Plasmid DNA from the normalized library NCI_CGAP
prepared, and ss circles were made in vitro. Fol
purification, this DNA was used as tracer in a s
hybridization reaction. The driver was PCR-ampli
from a pool of 5,000 clones made from the same l
(clones) 132376-132391, 1456007-1456775, and
1500552-1502855). Subtraction by Bento Soares an
Fatima Bonaldo."

ORIGIN
Query Match 23.7%; Score 326; DB 10; Length 337;
Best Local Similarity 100.0%; Pred.No. 1.2e-154;
Matches 326; Conservative 0; Mismatches 0; Indels 0;

Qy 1048 TCCCCACGGCCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCCCTTTGAGG
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Db 337 TCCCCACGGCCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCCCTTTGAGG
|||||


```
TCGACTCCCTCCCTGGCCACAGACACCCCGAGGCAATGTGTCACTGTACTCTGT 1167
TCGACTCCCTCCCTGGCCACAGACACCCCGAGGCAATGTGTCACTGTACTCTGT 245
AGATGGTCCAGAAAGACCCCACTTCAGGCACTTAAGAGGGGCTGGACCTGGCGGC 1227
AGATGGTCCAGAAAGACCCCACTTCAGGCACTTAAGAGGGGCTGGACCTGGCGGC 185
CCAAAGAGACTGGGCTAGCCAGGAGTCCCAAAATGTGAGGGGCGAGAAACAA 1287
CCAAAGAGACTGGGCTAGCCAGGAGTCCCAAAATGTGAGGGGCGAGAAACAA 125
CTCTCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATATATTTTATTAT 1347
CTCTCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATATATTTTATTAT 65
TGACAAATGTTGATAAATGG 1373
TGACAAATGTTGATAAATGG 39

1 .x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2297856 3',
sequence.
1.1 GI:5633396
piens (human)
piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 399)
p http://www.ncbi.nlm.nih.gov/ncicgap.
1 Cancer Institute, Cancer Genome Anatomy Project (CGAP),
ene Index
shed (1997)
: Robert Strausberg, Ph.D.
cgaps-r@mail.nih.gov
Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Buck, M.D., Ph.D.
library Preparation: M. Bento Soares, Ph.D.
library Arrayed by: Greg Lennon, Ph.D.
quencing by: Washington University Genome Sequencing Center
distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
.llnl.gov/bbrp/image/image.html
mer: -40UP from Gibco.
Location/Qualifiers
1. .399
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/clone="IMAGE:2297856"
/lab_host="DH10B"
/clone_lib="NCI CGAP Kid11"
/note="Organ: Kidney; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site 1: Not 1; Site 2: Eco RI;
Plasmid DNA from the normalized library NCI_CGAP Kid1 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneIDs 1322376-1323911, 1456007-1456775, and
1500552-1502855). Subtraction by Bento Soares and M.
Fatima Bonaldo. "
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23.6%; Score 324; DB 9; Length 399;

rity 100.0%; Pred. No. 1.3e-153; Indels 0; Gaps 0;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 347 CCCACCGCCCACTCTCCACCTCACTAGCTCCCAATCCCTTGAGGCCCC
QY 1110 TGATCTCGACTCCCTCCCTGGCCACAGACCCCGAGGCAATGTGTCACTGTACTCTG
DB 287 TGATCTCGACTCCCTCCCTGGCCACAGACCCCGAGGCAATGTGTCACTGTACTCTG
QY 1170 GCAAGGATGGTCCAGAAAGACCCCACTTCAGGCACTTAAGAGGGGCTGGACCTGGCGG
DB 227 GCAAGGATGGTCCAGAAAGACCCCACTTCAGGCACTTAAGAGGGGCTGGACCTGGCGG
QY 1230 GAAGCCAAAGAGACTGGGCTAGCCAGGAGTCCCAAAATGTGAGGGGCGAGAAACA
DB 167 GAAGCCAAAGAGACTGGGCTAGCCAGGAGTCCCAAAATGTGAGGGGCGAGAAACA
QY 1290 CAAGCTCTCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATATTTTATTAT
DB 107 CAAGCTCTCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATATTTTATTAT
QY 1350 TTGTGACAAATGTTGATAAATGG 1373
DB 47 TTGTGACAAATGTTGATAAATGG 24
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RESULT 35

BI966255

LOCUS

DEFINITION

BI966255

ACCESION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

1. .456

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:5672623"

/sex="Both"

/tissue_type="Islets of Langerhans"

/dev_stage="Adult"

/lab_host="DH10B"

/clone_lib="Melton Normalized Human Islet 4 N4-HIS

rough the I.M.A.G.E. Consortium/LLNL at:

image.llnl.gov

LLAM12814 row: n column: 16

ality sequence start: 74

ality sequence stop: 420.

Location/Qualifiers

1. .1819

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/clone_lib="NIH MGC 114"

/note="Organ: Brain; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA source anonymous pool of 6 male brains, age range 23-27 yo. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.5 kb, insert size range 1-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 019. Note: this is a NIH_MGC Library."

23.4%; Score 321; DB 12; Length 1819;

ality 100.0%; Pred. No. 5.9e-152;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

ICCACTCTCCACTCAGTCTCCCAATCCCTGACCCCTTTGAGGCCCCCAGTGA 1112

ICCACTCTCCACTCAGTCTCCCAATCCCTGACCCCTTTGAGGCCCCCAGTGA 139

ACTCCCCCTGGCCACAGCCCCCAGGGGCTGTCTCACTGTACTCTGTGGCA 1172

ACTCCCCCTGGCCACAGCCCCCAGGGGCTGTCTCACTGTACTCTGTGGCA 199

GGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGGCTGGACCTGGCGGAGAA 1232

GGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGGCTGGACCTGGCGGAGAA 259

AGAGACTGGGCTAGGCGAGGATCCCAATGTGAGGGGGCGAGAACAGACAA 1292

AGAGACTGGGCTAGGCGAGGATCCCAATGTGAGGGGGCGAGAACAGACAA 319

TCCCTTGAGAAATCCCTGTGGATTTTAAACAGATATTATTTTATTATTG 1352

TCCCTTGAGAAATCCCTGTGGATTTTAAACAGATATTATTTTATTATTG 379

AAATGTTGATAAATGG 1373

AAATGTTGATAAATGG 400

19 374 bp mRNA linear EST 14-FEB-2002

5.x1 Human insulinoma Homo sapiens cDNA 3', mRNA sequence.

19 1 GI:18669065

apiens (human)

apiens

ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

ses 1 to 374)

, D., Brown, J., Kenty, G., Permutt, A., Lee, C., Kaestner, K.,

ka, I., Searce, M., Brestelli, J., Gradwohl, G., Clifton, S.,

r, L., Marra, M., Pape, D., Wylie, T., Martin, J., Blistain, A.,

t, A., Theising, B., Ritter, E., Ronko, I., Bennett, J.,

as, M., Gibbons, M., McCann, R., Cole, R., Tsagareishvili, R.,

ms, T., Jackson, Y. and Bowers, Y.

ine Pancreas Consortium

ished (2000)

COMMENT

Other ESTs: ihl5b05.y1
Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
Endocrine Pancreas Consortium
Harvard University, Howard Hughes Medical Institute
Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cam
MA 02138

Tel: 617-495-1812

Fax: 617-495-8557

Email: dmelton@biohp.harvard.edu

Library was constructed by Dr. J. Ferrer In vivo mass-excis
pBluescript SK- by Dr. H. Inoue DNA sequencing by: Washing
University Genome Sequencing Center For information on obtai
clone please contact: Dr. Hiroshi Inoue (hinoue@im.wustl.ec
Seq primer: -40UP from Gibco.

FEATURES

source

1. .374

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/tissue_type="insulinoma"

/lab_host="DH10B (phage-resistant)"

/clone_lib="Human insulinoma"

/note="Organ: pancreas; Vector: pBluescript SK-; (xhoI; Site 2: EcoRI; Constructed with lambda ZAPII (Stratagene) by Dr. J. Ferrer, in vivo mass-excis pBluescript SK- by Dr. H. Inoue following the War University protocol (http://genome.wustl.edu/est/lambda_protocol.shtml. Please contact Hiroshi Inoue, MD/PhD for further information on this library (Metabolism Division Laboratory, Washington University School of Medicine, 8127, 660 S Euclid Ave, St. Louis, MO 63110). Note: is a Washington University Pancreas EST project 1;

ORIGIN

Query Match 23.3%; Score 320; DB 12; Length 374;
Best Local Similarity 100.0%; Pred. No. 1.4e-151; Indels 0; G
Matches 320; Conservative 0; Mismatches 0;

QY 1054 ACCGCCCACTCTCCACTCAGTCTCCCAATCCCTGACCCCTTTGAGGCCCCCAG

Db 346 ACCGCCCACTCTCCACTCAGTCTCCCAATCCCTGACCCCTTTGAGGCCCCCAG

QY 1114 CTCGACTCCCCCTGGCCACAGACCCCGGCGCATTTGTTCTACTGTCTGTGG

Db 286 CTCGACTCCCCCTGGCCACAGACCCCGGCGCATTTGTTCTACTGTCTGTGG

QY 1174 GATGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGCTCGACTGCGGCGAG

Db 226 GATGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGGCTCGACTGCGGCGAG

QY 1234 CCAAGAGACTGGGCTTAGGCCAGGAGTTCCTCAATCTGAGGGCGGAGAACAGA

Db 166 CCAAGAGACTGGGCTTAGGCCAGGAGTTCCTCAATCTGAGGGCGGAGAACAGA

QY 1294 CTCCTCCCTTGAGAAATCCCTGTGGATTTTAAACAGATATTATTTTATTATTA

Db 106 CTCCTCCCTTGAGAAATCCCTGTGGATTTTAAACAGATATTATTTTATTATTA

QY 1354 GACAAATGTTGATAAATGG 1373

Db 46 GACAAATGTTGATAAATGG 27

RESULT 39

AI695776/c

LOCUS

DEFINITION

AI695776

AI695776

VERSION

KEYWORDS

SOURCE

AI695776 329 bp mRNA linear EST 17

wb77907.x1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:231

mRNA sequence.

AI695776

AI695776.1 GI:4983676

EST.

Homo sapiens (human)

apiens
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ses 1 to 329)
AP http://www.ncbi.nlm.nih.gov/ncicgap.
al Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Gene Index
ished (1997)
t: Robert Strausberg, Ph.D.
cgapbs-remail.nih.gov
Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R.
-Buck, M.D., Ph.D.
Library Preparation: M. Bento Soares, Ph.D.
Library Arrayed by: Greg Lennon, Ph.D.
sequencing by: Washington University Genome Sequencing Center
distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
o.llnl.gov/bbrp/image/image.html
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Location/Qualifiers
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/note="Organ: prostate; Vector: p7T3D-Pac (Pharmacia)
with a modified polylinker; plasmid DNA from the
normalized library NCI CGAP Pr22 was prepared, and as
circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones from the same library (cloneids
985608-986759, 1101192-1101959, and 1217928-1220615).
Subtraction by Bento Soares and M. Fatima Bonaldo."

23.2%; Score 318; DB 9; Length 329;
arity 100.0%; Pred. No. 1.4e-150;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
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CACTCTCCACCTCACTAGTCCGCAATCCCTGACCCCTTGAGGCCCCAGTGATCT 270
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TCCCCCTGGCCACAGACCCCCAGGGCATTGTGTTCACTGTACTCTGTGGCAAGG 1235
TCCCCCTGGCCACAGACCCCCAGGGCATTGTGTTCACTGTACTCTGTGGCAAGG 150
TCCCCCTGGCCACAGACCCCCAGGGCATTGTGTTCACTGTACTCTGTGGCAAGG 1295
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CCCTTGAGAAATTCCTGTGGATTTTAAACAGATATTTATTTATTTATTTATTTGTA 1355
CCCTTGAGAAATTCCTGTGGATTTTAAACAGATATTTATTTATTTATTTATTTGTA 30
AATGTTGATAAATGG 1373
AATGTTGATAAATGG 12

LOCUS BF195436 340 bp mRNA linear EST 03
DEFINITION 7n17g12.x1 NCI_CGAP_Brn23 Homo sapiens cDNA clone IMAGE:35
mRNA sequence.
ACCESSION BF195436
VERSION BF195436.1 GI:11082306
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 340)
AUTHORS NCI/NINDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute / National Institute of Neurolog
Disorders and Stroke, Brain Tumor Genome Anatomy Project
(CGAP/BTGAP), Tumor Gene Index
Unpublished (1998)
JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgapbs-remail.nih.gov
Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenfe
Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Pati
Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencin
Clone distribution: NCI-CGAP clone distribution informati
found through the I.M.A.G.E. Consortium/LLNL, send email t
info@image.llnl.gov.
FEATURES
Location/Qualifiers
1. 340
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3564887"
/tissue_type="glioblastoma (pooled)"
/lab_host="DH10B"
/clone_lib="NCI CGAP Brn23"
/note="Organ: brain; Vector: p7T3D-Pac (Pharmac
modified polylinker; Site 1: Not I; Site 2: Eco I
strand cDNA was primed with a Not I - oligo(dT) I
TGTACCAATCTGAAGTGGAGCGCGGCATATCTTTTTTTTTTTT
T 3'; double-stranded cDNA was ligated to Eco I
adaptors (Pharmacia), digested with Not I and C
the Not I and Eco RI sites of the modified p7T
Library is normalized, and was constructed by B
Soares and M. Fatima Bonaldo."

ORIGIN
Query Match 23.2%; Score 318; DB 10; Length 340;
Best Local Similarity 100.0%; Pred. No. 1.4e-150;
Matches 318; Conservative 0; Mismatches 0; Indels 0;
QY 1056 CGCCCACTCTCCACCTCACTAGTCCGCAATCCCTTGAGGCCCCAGT
DB 340 CGCCCACTCTCCACCTCACTAGTCCGCAATCCCTTGAGGCCCCAGT
QY 1116 CGACTCCCCCTGGCCACAGACCCCGAGGGCATTGTGTTCACTGTA
DB 280 CGACTCCCCCTGGCCACAGACCCCGAGGGCATTGTGTTCACTGTA
QY 1176 ATGGGTCCAGAGAGACCCCACTTCAGGCACCTAAGAGGGGCTGGAGCCGAGG
DB 220 ATGGGTCCAGAGAGACCCCACTTCAGGCACCTAAGAGGGGCTGGAGCCGAGG
QY 1236 AAGAGAGACTGGGCCCTAGGCCAGAGTTCCTCAATGTGAGGGGCGAGAAACAGAC
DB 160 AAGAGAGACTGGGCCCTAGGCCAGAGTTCCTCAATGTGAGGGGCGAGAAACAGAC
QY 1296 CCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATTTATTTATTTAT
DB 100 CCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATTTATTTATTTAT
QY 1356 CAAATGTTGATAAATGG 1373

IGTTGATAAATGG 23

5 .xl Soares_NFL_T_GBC_S1 mRNA linear EST 30-NOV-1998
842906 3', mRNA sequence.

5.1 GI:3804188

piens (human)

piens

a; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

a; Eutheria; Primates; Catarrhini; Hominidae; Homo.

es 1 to 407

P <http://www.ncbi.nlm.nih.gov/ncicgap>.

1 Cancer Institute, Cancer Genome Anatomy Project (CGAP),

ene Index

shed (1997)

: Robert Strausberg, Ph.D.

cgapbs-r@mail.nih.gov

one is available royalty-free through LLNL; contact the

onsortium (info@image.llnl.gov) for further information.

Length: 610 Std Error: 0.00

mer: -40UP from Gibco

ality sequence stop: 401.

Location/Qualifiers

1. .407

/organism="Homo sapiens"

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/notes="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with

a modified polylinker; Site 1: Not I; Site 2: Eco RI;

Equal amounts of plasmid DNA from three normalized

libraries (fetal lung NbHL19W, testis NHT, and B-cell

NCI CGAP GCBI) were mixed and ss circles were made in

vitro. Following HAP purification, this DNA was used as

tracer in a subtractive hybridization reaction. The driver

was PCR-amplified cDNAs from pools of 5,000 clones made

from the same 3 libraries. The pools consisted of

I.M.A.G.E. clones 297480-302087, 682632-687239,

726408-728711, and 729096-731399. Subtraction by Bento

Soares and M. Fatima Bonaldo.

23.0%; Score 316; DB 9; Length 407;

urity 100.0%; Pred. No. 1.5e-149;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

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TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 276

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 1177

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 216

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 1237

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 156

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 1297

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 96

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 1357

TCTCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCG 36

QY 1358 AAATGTTGATAAATGG 1373

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Db 35 AAATGTTGATAAATGG 20

RESULT 42

AI291866/c

LOCUS

DEFINITION

qmb6c02.x1 NCI_CGAP_Lu5 Homo sapiens cDNA clone IMAGE:18956

mRNA sequence.

ACCESSION

AI291866

VERSION

AI291866.1 GI:3934640

KEYWORDS

EST.

SOURCE

Homo sapiens (human)

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutelec

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 416)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (C

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Mici

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing

Clone distribution: NCI-CGAP clone distribution informati

found through the I.M.A.G.E. Consortium/LLNL at:

www.bio.llnl.gov/bbrp/image/image.html

Insert Length: 649 Std Error: 0.00

Seg primer: -40UP from Gibco

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Location/Qualifiers

1. .416

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/clone_lib="NCI_CGAP_Lu5"

/notes="Organ: lung; Vector: pT7T3D-Pac (Pharmacia

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Not I - oligo(dT) primer. Double-stranded cDNA wa

to Eco RI adaptors (Pharmacia), digested with Not

cloned into the Not I and Eco RI sites of the mod

pT7T3 vector. Library is normalized. Library was

constructed by Bento Soares and M. Fatima Bonaldo

Query Match 22.5%; Score 309; DB 9; Length 416;

Best Local Similarity 100.0%; Pred. No. 5.7e-146; Indels 0; G

Matches 309; Conservative 0; Mismatches 0; Indels 0; G

QY 1065 TCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCGACT

|||||

Db 334 TCCACCTCCTAGTCTCCCAATCCCTGACCCCTTGAGGCCCCAGTATCTCGACT

|||||

QY 1125 CTGTGGCCACAGACCCCGAGGGCATTTGTGTTTCACTGTACTGTGGGGCAAGATGGG

|||||

Db 274 CTGTGGCCACAGACCCCGAGGGCATTTGTGTTTCACTGTACTGTGGGGCAAGATGGG

|||||

QY 1185 GAAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGCGGCGAGGAAGCCAAAGA

|||||

Db 214 GAAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGCGGCGAGGAAGCCAAAGA

|||||

QY 1245 GGGCCTAGGGCAGGAGTTCCCAATGTAGGGGCGAGAAACAGCAAGCTCTCTCC

|||||

Db 154 GGGCCTAGGGCAGGAGTTCCCAATGTAGGGGCGAGAAACAGCAAGCTCTCTCC


```

mer: M13 FORWARD
es.
Location/Qualifiers
1. 333
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clones="UI-E-CL1-afe-m-20-0-UI"
/tissue_type="human retina"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/clone_lib="UI-E-CL1"
/notes="Organ: eye; Vector: pT73-Pac (Pharmacia) with a
modified polylinker; Site 1: EcoR I; Site 2: Not I;
UI-E-CL1 is a normalized cDNA library containing the
following tissue(s): retina. The library was constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT73-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tag for this library is CCGCG. This library was
created for the program, Gene Discovery in the Visual
System, supported by National Eye Institute (NEI).
TAG TISSUE=human retina
TAG_LIB=UI-E-CL1
TAG_SEQ=CCGCG"

21.6%; Score 296; DB 13; Length 333;
arity 100.0%; Pred. No. 2.4e-139; Indels 0; Gaps 0;
onservative 0; Mismatches 0;

CCAAATCCCTGACCTTTGAGGCGCCCGCCAGTATCTCGACTCCCTCCCTGGCCACAGAC 1137
CCAAATCCCTGACCTTTGAGGCGCCCGCCAGTATCTCGACTCCCTCCCTGGCCACAGAC 274
AGGGCATTTGTGTTACTGTTCTGTGGGCAAGGATGGTCCAGAAAGCCCACTT 1197
AGGGCATTTGTGTTACTGTTCTGTGGGCAAGGATGGTCCAGAAAGCCCACTT 214
CACTAAGAGGGCTGAGCTGGCGGCGAGCAAGCCAAAGAGACTGGCCCTAGGCCAG 1257
CACTAAGAGGGCTGAGCTGGCGGCGAGCAAGCCAAAGAGACTGGCCCTAGGCCAG 154
TCCCAAAATGTGAGGGGCGAGAAACAAGACAGCTCCTCCCTTGAGAAATCCCTGTG 1317
TCCCAAAATGTGAGGGGCGAGAAACAAGACAGCTCCTCCCTTGAGAAATCCCTGTG 94
TTTAAACAGATATTATTTTATTTATTTATTTATTTGACAAATGTTGATAATGG 1373
TTTAAACAGATATTATTTTATTTATTTATTTATTTGACAAATGTTGATAATGG 38

146 298 bp mRNA linear EST 28-FEB-2002
Q1-acs-a-05-0-UI.r1 UI-E-CQ1 Homo sapiens cDNA clone
Q1-acs-a-05-0-UI 5', mRNA sequence.
146
146.1 GI:19002204
sapiens (human)
sapiens
yota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
lia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ases 1 to 298)
do.M.F., Lennon,G. and Soares,M.B.
lization and subtraction: two approaches to facilitate gene
very

```

```

JOURNAL Genome Res. 6 (9), 791-806 (1996)
MEDLINE 97044477
PUBMED 8889548
COMMENT Contact: Soares, MB
Coordinated Laboratory for Computational Genomics
University of Iowa
375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: bento-soares@uiowa.edu
Tissue Procurement: Dr. Gregg Hageman
cDNA Library preparation: Dr. M. Bento Soares, University
cDNA Library Arrayed by: Dr. M. Bento Soares, University
DNA sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Researchers may obtain clones from Re
Genetics (www.resgen.com).
Seq primer: M13 Reverse.
Location/Qualifiers
1. 298
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clones="UI-E-CQ1-acs-a-05-0-UI"
/tissue_type="optic nerve"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies) (T1 phage re
/clone_lib="UI-E-CQ1"
/notes="Organ: eye; Vector: pT73-Pac (Pharmacia)
modified polylinker; Site 1: EcoR I; Site 2: Not
UI-E-CQ1 is a normalized cDNA library containing
following tissue(s): optic nerve. The library was
constructed according to Bonaldo, Lennon and Soar
Genome Research, 6:791-806, 1996. First strand c
synthesis was primed with an oligo-dT primer cont
Not I site. Double stranded cDNA was ligated to a
adaptor, digested with Not I, and cloned directic
into pT73-Pac vector. The oligonucleotide used t
the synthesis of first-strand cDNA contains a lib
sequence that is located between the Not I site a
(dT)18 tail. The sequence tag for this library is
CCATTAAAGT. This library was created for the prog
Discovery in the Visual System, supported by Nati
Institute (NEI)."

```

ORIGIN

```

Query Match 21.1%; Score 290; DB 12; Length 298;
Best Local Similarity 100.0%; Pred. No. 2.7e-136;
Matches 290; Conservative 0; Mismatches 0; Indels 0; C
QY 462 GGACGGAGCGCAGGAGCTGTGGACGGGACAGTGGCTGGGAGGAGCCAGAG
DB 9 GGACGGAGCGCAGGAGCTGTGGACGGGACAGTGGCTGGGAGGAGCCAGAG
QY 522 CAGCTCCAGCCCTCTGCGCTACACCGCCAGATCGGGAGGTTTATGTCACCCCGK
DB 69 CAGCTCCAGCCCTCTGCGCTACACCGCCAGATCGGGAGGTTTATGTCACCCCGK
QY 582 GCTCTACTACTGTACTGTCTAGGTCACATTTGATGAGGGAGGCTGTCTACTGT
DB 129 GCTCTACTACTGTACTGTCTAGGTCACATTTGATGAGGGAGGCTGTCTACTGT
QY 642 GGACTTGTGTTGGTGGTGTGCTGGGCGCCCTGGCTGGAGGAAATTTCTCAGCCJ
DB 189 GGACTTGTGTTGGTGGTGTGCTGGGCGCCCTGGCTGGAGGAAATTTCTCAGCCJ
QY 702 GGCCAGTTTCCCTCGGGGCGCCAGCTCCGCTCTGCCAGGTTGTCTGGGCTGT 751
DB 249 GGCCAGTTTCCCTCGGGGCGCCAGCTCCGCTCTGCCAGGTTGTCTGGGCTGT 298

```

RESULT 48
BG686319
LOCUS

587 bp mRNA linear EST 0

32F1 NIH_MGC_48 Homo sapiens cDNA clone IMAGE:4766071 5',
sequence.

9
9.1 GI:13917716

piens (human)

Diens

ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.

es 1 to 587)

http://imgc.nci.nih.gov/.

1 Institutes of Health, Mammalian Gene Collection (MGC)

shed (1999)

: Robert Strausberg, Ph.D.

cgapbs@mail.nih.gov

Procurement: Louis M. Staudt, M.D., Ph.D.

Library Preparation: Ling Hong/Rubin Laboratory

Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

quencing by: Incyte Genomics, Inc.

distribution: MGC clone distribution information can be

through the I.M.A.G.E. Consortium/LLNL at:

image.llnl.gov

LLCM1625 row: p column: 08

ality sequence stop: 587.

Location/Qualifiers

1..587

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:4766071"

/tissue_type="primary B-cells from tonsils (cell line)"

/lab_host="DH10B (phage-resistant)"

/clone_lib="NIH_MGC_48"

/notes="Organ: B-cells; Vector: pOTB7; Site: 1: XhoI;

Site 2: EcoRI; cDNA made by oligo-dT priming.

Directionally cloned into EcoRI/XhoI sites using the

following 5' adaptor: GGCACGAG(G). Size-selected >500bp

for average insert size 1.8kb. Library constructed by Ling

Hong in the laboratory of Gerald M. Rubin (University of

California, Berkeley) using ZAP-cDNA synthesis kit

(Stratagene) and Superscript II RT (Life Technologies).

Note: this is a NIH_MGC Library."

20.6%; Score 283; DB 12; Length 587;

urity 99.6%; Pred. No. 1.2e-132;

nservative 0; Mismatches 1; Indels 1; Gaps 1;

XCGCGCGGTCCCGCTCCCGGATCCCTCGGGTCCCGGATGGGGGGGGGTGAGG 84

XCGCGCGGTCCCGCTCCCGGATCCCTCGGGTCCCGGATGGGGGGGGGTGAGG 76

XACAGCCCCCGCCCGATGGCCCGCTCGGAGCCAGAGCGGAGGGGGCGCGG 144

XACAGCCCCCGCCCGATGGCCCGCTCGGAGCCAGAGCGGAGGGGGCGCGG 136

AGCGGGCACCAGCCCTGCTGGTCCCGCTCGGAGCCAGAGCGGAGGGGGCGCGG 203

AGCGGGCACCAGCCCTGCTGGTCCCGCTCGGAGCCAGAGCGGAGGGGGCGCGG 196

CTCGGCTCTGCTGGCGGTGGTTCAGTTTGGGAGCGGGGATCGGTCGCCGCCA 263

CTCGGCTCTGCTGGCGGTGGTTCAGTTTGGGAGCGGGGATCGGTCGCCGCCA 256

CCTGCCAGGAGGAGTGGTGGCAGAGGAGGACAGGACCCGTCGGAATGCC 323

CCTGCCAGGAGGAGTGGTGGCAGAGGAGGACAGGACCCGTCGGAATGCC 316

ACAGAGAGAGGAGGAGTCTGCGCCCTTCTGACCGACTAGTTGGGCTCCGAG 383

ACAGAGAGAGGAGGAGTCTGCGCCCTTCTGACCGACTAGTTGGGCTCCGAG 376

GCACCTAAAGCGCGGAAACACAGGGCTCGAAGAGCGGATCGAGCCCATTTAAGT 443

Db 377 AAGTGCACCTAAAGCGCGGAAACACAGGGCTCGAAGAGCGATCGAGCCCATTTATGA

Qy 444 TCATCCAGCAGCTCGACAGAGCGGAGCGGAGCGAG 478

Db 437 TCATCCAGCAGCTCGACAGAGCGGAGCGGAGCGAG 471

RESULT 49

AF163779

LOCUS

DEFINITION

AF163779 Human Homo sapiens genomic clone BAC750E14, genom

sequence.

ACCESSION

AF163779

VERSION

AF163779.1 GI:5726439

KEYWORDS

GSS.

SOURCE

Homo sapiens (human)

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutelec

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 1027)

Authors

Cousin, P., Billotte, J., Chaubert, P. and Shaw, P.H.

Physical map of 17p13 and the genes adjacent to p53

JOURNAL

Genomics 63 (1), 60-68 (2000)

MEDLINE

20130114

PUBMED

10662545

COMMENT

Contact: Shaw PH

Experimental Oncology

Institute of Pathology

Rue du Bugnon 25, Lausanne, VD 1011, Switzerland

sub_clone=AB2R Asc-BamHI PSL1160

Classes: BAC subclone.

Location/Qualifiers

1..1027

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/map="17p"

/clone="BAC750E14"

/clone_lib="Human"

ORIGIN

Query Match 19.7%; Score 271; DB 28; Length 1027;

Best Local Similarity 99.7%; Pred. No. 1.7e-126;

Matches 321; Conservative 0; Mismatches 1; Indels 0; G

Qy 1052 CCACCCCGCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCCCTTTAGGCCCC

Db 268 CCACCCCGCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCCCTTTAGGCCCC

Qy 1112 ATCTCGACTCCCGCTGGCCACAGACCCCGAGGCGATTGTTCACGTACTCTGT

Db 328 ATCTCGACTCCCGCTGGCCACAGACCCCGAGGCGATTGTTCACGTACTCTGT

Qy 1172 AAGGATGGGTTCAGAGAGACCCCACTTCAGGCACTAAGAGGGCTGACCTGGCGGC

Db 388 AAGGATGGGTTCAGAGAGACCCCACTTCAGGCACTAAGAGGGCTGACCTGGCGGC

Qy 1232 AGCCAAAGAGACTGGGCTAGGCGGAGGTTCCCAATGTGAGGGCGAGAAACAA

Db 448 AGCCAAAGAGACTGGGCTAGGCGGAGGTTCCCAATGTGAGGGCGAGAAACAA

Qy 1292 AGCTCTCCCTTCAGAAATTCCTGTGGATTTTAAAAACAGATATTTATTTAT

Db 508 AGCTCTCCCTTCAGAAATTCCTGTGGATTTTAAAAACAGATATTTATTTAT

Qy 1352 GTGACAAAATGTTGATAAATGG 1373

Db 568 GTGACAAAATGTTGATAAATGG 589

RESULT 50

AI760777/c

77 5.xl NCI_CGAP_Kid12 346 bp mRNA linear EST 24-JUN-1999
sequence. IMAGE:2398377 3',
77
77.1 GI:5176444
apiens (human)
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
ses 1 to 346)
AP http://www.ncbi.nlm.nih.gov/ncicgap.
al Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Gene Index
ished (1997)
t: Robert Strausberg, Ph.D.
cgapbs-remail.nih.gov
Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
-Buck, M.D., Ph.D.
Library Preparation: M. Bento Soares, Ph.D.
Library Arrayed by: Greg Lennon, Ph.D.
sequencing by: Washington University Genome Sequencing Center
distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
o.llnl.gov/bbrp/image/image.html
imer: -40UP from Gibco.
Location/Qualifiers
1..346
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2398377"
/tissue_type="2 pooled tumors (clear cell type)"
/lab_host="DH10B"
/clone_lib="NCI_CGAP_Kid12"
/note="Organ: Kidney; Vector: pT73D-Pac (Pharmacia) with
a modified polylinker; Site 1: Not 1; Site 2: Eco RI;
Plasmid DNA from the normalized library NCI CGAP Kid5 was
prepared, and ss circles were made in vitro following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneIDs 1323912-1325831, 1471368-1472903 and
1492104-1493255). Subtraction by Bento Soares and M.
Fatima Bonaldo."
19.4%; Score 267; DB 9; Length 346;
arity 99.7%; Pred. No. 1.5e-124;
onservative 0; Mismatches 1; Indels 0; Gaps 0;
CAGCTCCACCTCAGTCTCCCATCCCTGACCTTTGAGGCCCCCAGTGATCT 1115
CAGCTCCACCTCAGTCTCCCATCCCTGACCTTTGAGGCCCCCAGTGATCT 287
CTCCCCCTGGCCACAGACCCAGGCGCATGTGTTTACCTGTGTTGGGCAAGG 1175
CTCCCCCTGGCCACAGACCCAGGCGCATGTGTTTACCTGTGTTGGGCAAGG 227
GGTCCAGAGACCCCACTTCCAGGCGCATGTGTTTACCTGTGTTGGGCAAGG 1235
GGTCCAGAGACCCCACTTCCAGGCGCATGTGTTTACCTGTGTTGGGCAAGG 167
GAGACTGGGCTAGCCAGGAGTTCCTCAATGTGAGGGGCGAGAAACAGCAAGCT 1295
GAGACTGGGCTAGCCAGGAGTTCCTCAATGTGAGGGGCGAGAAACAGCAAGCT 107
CCCTTGAGAAATTCCTGTGATTTTAAACAGATATTTATTTATTTATTTGTA 1355
CCCTTGAGAAATTCCTGTGATTTTAAACAGATATTTATTTATTTATTTGTA 47
AATGTTGATAATGG 1373

Db 46 CAAAATGTTGATAAATGG 29
RESULT 51
BF940141/c
LOCUS
DEFINITION
BF940141 346 bp mRNA linear EST 22
nac68g06.x1 NCI_CGAP_Brn23 Homo sapiens cDNA clone IMAGE:3
similar to contains Element MSRI repetitive element ;, MRN
sequence.
ACCESSION
BF940141
VERSION
BF940141.1 GI:12357461
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
1 (bases 1 to 346)
NCI/NINDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute / National Institute of Neurolog
Disorders and Stroke, Brain Tumor Genome Anatomy Project
(CGAP/BTGP), Tumor Gene Index
Unpublished (1998)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenfe
Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fati
Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencin
found through the I.M.A.G.E. Consortium/LLNL, send email t
info@image.llnl.gov
Seq primer: -40UP from Gibco
High quality sequence stop: 321.
Location/Qualifiers
1..346
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3439667"
/tissue_type="glioblastoma (pooled)"
/lab_host="DH10B"
/clone_lib="NCI_CGAP_Brn23"
/note="Organ: brain; Vector: pT73D-Pac (Pharmac
modified polylinker; Site 1: Not 1; Site 2: Eco I
strand cDNA was primed with a Not I - oligo(dT) I
TGTTACCAATCTGAAGTGGGAGCGCGCATATCTTTTTTTTTT
T 3'; double-stranded cDNA was ligated to Eco I
adaptors (Pharmacia), digested with Not I and c
the Not I and Eco RI sites of the modified pT7T
Library is normalized, and was constructed by B
Soares and M.Fatima Bonaldo."
Query Match 19.1%; Score 262; DB 10; Length 346;
Best Local Similarity 100.0%; Pred. No. 5.4e-122; Indels 0;
Matches 262; Conservative 0; Mismatches 0;
QY 1048 TCCCCCAGCGCCACTCTCCACTCAGTCTCCCAATCCCTGACCCCTTTGAGG
346 TCCCCCAGCGCCACTCTCTCCACTCAGTCTCCCAATCCCTGACCCCTTTGAGG
1108 AGTGATCTCGACTCCCTCCCTGGCCACAGACCCCGGCGCATTTGTTCTACTGTAC
286 AGTGATCTCGACTCCCTCCCTGGCCACAGACCCCGGCGCATTTGTTCTACTGTAC
1168 GGGCAAGGATGGGTTCAGAAAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTG
226 GGGCAAGGATGGGTTCAGAAAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTG
1228 AGGAAGCCAAAGAGACTGGGCTAGCCAGGAGTTCCTCAATGTGAGGGGCGGAGA

3CCAGAGAGACTGGGCGCTAGCCAGGAGTTCCTCCAAATGTGAGGGCGGAGAAACAA 107

3CTCCTCCCTTGAGAAAT 1309

3CTCCTCCCTTGAGAAAT 85

8 561 bp mRNA linear EST 25-JAN-2001
2.x1 NCI CGAP Brn23 Homo sapiens cDNA clone IMAGE:3441742 3'
to TR:Q9UK76 Q9UK76 HN1 PROTEIN. ; mRNA sequence.

8.1 GI:12512043

piens (human)

piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.

es 1 to 561)

IDS-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

rs and Stroke, Brain Tumor Genome Anatomy Project

TCAP), Tumor Gene Index

shed (1998)

: Robert Strausberg, Ph.D.

cgapbs-r@mail.nih.gov

Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D.,

library Preparation: M. Bento Soares, Ph.D., M. Fatima

, Ph.D.

quencing by: Greg Lennon, Ph.D.

distribution: NCI-CGAP clone distribution information can be

through the I.M.A.G.E. Consortium/LLNL, send email to:

age.llnl.gov

ality sequence stop: 260.

Location/Qualifiers

1. 561

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:3441742"

/tissue_type="glioblastoma (pooled)"

/lab_host="DH10B"

/clone_lib="NCI CGAP Brn23"

/note="Organ: brain; Vector: pT73D-Pac (Pharmacia) with a

modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st

strand cDNA was primed with a Not I - oligo(dT) primer [5'

TGTTACCAATCTGAAGTGGAGCGCGCATATCTTTTTTTTTTTTTTTTTTT

T 3']; double-stranded cDNA was ligated to Eco RI

adaptors (Pharmacia), digested with Not I and cloned into

the Not I and Eco RI sites of the modified pT73 vector.

Library is normalized, and was constructed by Bento

Soares and M. Fatima Bonaldo."

18.0%; Score 247; DB 10; Length 561;

arity 100.0%; Pred. No. 2.7e-114;

onservative 0; Mismatches 0; Indels 0; Gaps 0;

CACAGACCCCGAGGCAATGTGTTCACTGTACTCTGTGGCGAGGATGGTCCAGA 1186

CACAGACCCCGAGGCAATGTGTTCACTGTACTCTGTGGCGAGGATGGTCCAGA 212

CCCACTTCAGGCACTAAGAGGGGCTGGACCTGGCGGAGGATGGGCTTAGG 1246

CCCACTTCAGGCACTAAGAGGGGCTGGACCTGGCGGAGGATGGGCTTAGG 152

AGGCCAGGAGTTCCTCCAAATGTGAGGGCGGAGAAACAAGACAGCTCCTCCCTGAG 1306

AGGCCAGGAGTTCCTCCAAATGTGAGGGCGGAGAAACAAGACAGCTCCTCCCTGAG 92

QY 1307 AATTCCTGTGGATTTTAAACAGATATTATTTTATTATTATTATTTGTGACAAATGT

Db 91 AATTCCTGTGGATTTTAAACAGATATTATTTTATTATTATTATTTGTGACAAATGT

QY 1367 TAAATGG 1373

Db 31 TAAATGG 25

RESULT 53

AW081731/c

LOCUS

DEFINITION

AW081731 318 bp mRNA linear EST 14-

xb70a02.x1 Soares NFL T GBC S1 Homo sapiens cDNA clone

IMAGE:2581610 3', similar to contains element MSRI repetitiv

element ; mRNA sequence.

AW081731 GI:6036883

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 318)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project ((

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

This clone is available royalty-free through LLNL ; contac

IMAGE Consortium (info@image.llnl.gov) for further inform

Seq primer: -40UP from Gibco

High quality sequence stop: 314.

Location/Qualifiers

1. 318

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:2581610"

/lab_host="DH10B"

/clone_lib="Soares NFL T GBC S1"

/note="Organ: pooled; Vector: pT73D-Pac (Pharmac

a modified polylinker; Site 1: Not I; Site 2: Eco

Equal amounts of plasmid DNA from three normalize

libraries (fetal lung MbHL19W, testis NHT, and B-

NCI CGAP GCBI) were mixed, and ss circles were ma

vitro. Following HAP purification, this DNA was u

tracer in a subtractive hybridization reaction. T

was PCR-amplified cDNAs from pools of 5,000 clone

from the same 3 libraries. The pools consisted of

I.M.A.G.E. clones 297480-302087, 682632-687239,

726408-728711, and 729096-731399. Subtraction by

Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 17.2%; Score 236; DB 9; Length 318;

Best Local Similarity 100.0%; Pred. No. 9.9e-109;

Matches 236; Conservative 0; Mismatches 0; Indels 0; G

QY 1138 CCCAGGCGCATTTGTTCACTGTACTCTGTGGCGAGGATGGGTCACAGAGACCCC

Db 254 CCCAGGCGCATTTGTTCACTGTACTCTGTGGCGAGGATGGGTCACAGAGACCCC

QY 1198 CAGCCTAAGAGGGGCTGGACCTGGCGGAGGATGGGTCACAGAGACTGGGCTAGG

Db 194 CAGCCTAAGAGGGGCTGGACCTGGCGGAGGATGGGTCACAGAGACTGGGCTAGG

QY 1258 GAGTTCCTCCAAATGTGAGGGCGGAGAAACAAGACAGCTCCTCCTTGAGATTCCC

Db 134 GAGTTCCTCCAAATGTGAGGGCGGAGAAACAAGACAGCTCCTCCTTGAGATTCCC

QY 1318 GATTTTAAACAGATATTATTTTATTATTATTATTGTGACAAATGTTGATAAATGG

```

|||||
TTTTAAACAGATATTATTTTATTTATTTATTTGTCGACAAAATGTTGATAAATGG 19
|||||

41 264 bp mRNA linear EST 06-APR-2000
0.xl Soares NFL T.GBC.S1 Homo sapiens cDNA clone
2978587 3', mRNA sequence.
41 41.1 GI:7454367
apiens (human)
apiens
ota: Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia: Eutheria; Primates; Catarrhini; Hominidae; Homo.
ues 1 to 264)
AP http://www.ncbi.nlm.nih.gov/ncicgap.
al Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Gene Index
ished (1997)
t: Robert Strausberg, Ph.D.
: cgapsb-r@mail.nih.gov
:one is available royalty-free through LNL; contact the
Consortium (info@image.llnl.gov) for further information.
imer: -40UP from Gibco
uality sequence stop: 263.
Location/Qualifiers
1..264
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2978587"
/lab_host="DH10B"
/clone_lib="Soares_NFL_T.GBC.S1"
/notes="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site 1: Not 1; Site 2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NbHL19W, testis NHT, and B-cell
NCL CGAP_GCB1) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo."
16.8%; Score 230; DB 10; Length 264;
larity 100.0%; Pred. No. 1.1e-105;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ATTGTGTTCACTGTACTCTGTGGCAAGGATGGTCCAGAGACCCCACTTCAGGCA 1203
ATTGTGTTCACTGTACTCTGTGGCAAGGATGGTCCAGAGACCCCACTTCAGGCA 195
AGAGGGGCTGGACCTGCGCGCAGAACCCAAAGAGACTGGGCCCTAGGCCAGGAGTTC 1263
AGAGGGGCTGGACCTGCGCGCAGAACCCAAAGAGACTGGGCCCTAGGCCAGGAGTTC 135
AATGTGAGGGGCGAGAAACAGACCAAGCTCTCCCTTTGAGAATTCCTCTGGATTTT 1323
AATGTGAGGGGCGAGAAACAGACCAAGCTCTCCCTTTGAGAATTCCTCTGGATTTT 75
AACAGATATTTATTTATTTATTTATTTGTCGACAAAATGTTGATAAATGG 1373
AACAGATATTTATTTATTTATTTATTTATTTGTCGACAAAATGTTGATAAATGG 25

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563 253 bp mRNA linear EST 21-DEC-1999

```

DEFINITION w17905.x1 NCI_CGAP Kid12 Homo sapiens cDNA clone IMAGE.24
similar to contains element PTR5 repetitive element ;, mRNA
sequence.
ACCESSION AF863563
VERSION AF863563.1 GI:5527670
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 253)
NCI_CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapsb-r@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Mic
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Sequencing by: Greg Lennon, Ph.D.
CDNA Library Arrayed by: Washington University Genome Sequencin
Clone distribution: NCI-CGAP clone distribution informati
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 669 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 253.
Location/Qualifiers
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/clone="IMAGE:2403128"
/tissue_type="2 pooled tumors (clear cell type)"
/lab_host="DH10B"
/clone_lib="NCI_CGAP_Kid12"
/notes="Organ: Kidney; Vector: pT7T3D-Pac (Pharma
a modified polylinker; Site 1: Not 1; Site 2: Eco
plasmid DNA from the normalized library NCI_CGAP
prepared, and ss circles were made in vitro. Fol
purification, this DNA was used as tracer in a s
hybridization reaction. The driver was PCR-ampli
from a pool of 5,000 clones made from the same 1
(cloneIDs 1323912-1325831, 1471368-1472903 and
1492104-1493255). Subtraction by Bento Soares an
Fatima Bonaldo."

```

ORIGIN

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Query Match 16.7%; Score 229; DB 9; Length 253;
Best Local Similarity 100.0%; Pred. No. 3.5e-105;
Matches 229; Conservative 0; Mismatches 0; Indels 0;

QY 1145 GCATTGTGTTCACTGTACTCTGTGGCAAGGATGGTCCAGAGACCCCACTTC
Db 253 GCATTGTGTTCACTGTACTCTGTGGCAAGGATGGTCCAGAGACCCCACTTC
QY 1205 TAAGAGGGGCTGGACCTGCGCGCAGGAGCCAAAGAGACTGGGCCCTAGGCCAGGA
Db 193 TAAGAGGGGCTGGACCTGCGCGCAGGAGCCAAAGAGACTGGGCCCTAGGCCAGGA
QY 1265 CAAATGTGAGGGGCGAGAAACAGACCAAGCTCTCCCTTTGAGAATTCCTCTGTGGA
Db 133 CAAATGTGAGGGGCGAGAAACAGACCAAGCTCTCCCTTTGAGAATTCCTCTGTGGA
QY 1325 AAAACAGATATTTATTTATTTATTTATTTGTCGACAAAATGTTGATAAATGG 1373
Db 73 AAAACAGATATTTATTTATTTATTTATTTGTCGACAAAATGTTGATAAATGG 25

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RESULT 56

AI682487/c 238 bp mRNA linear EST 1
LOCUS AI682487

apiens
ota: Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ses 1 to 351)
NDS-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
al Cancer Institute / National Institute of Neurological
ers and Stroke, Brain Tumor Genome Anatomy Project
BTGAP), Tumor Gene Index
ished (1998)
t: Robert Strausberg, Ph.D.
cgaps-remail.nih.gov
Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D.,
Library Preparation: M. Bento Soares, Ph.D., M. Fatima
o, Ph.D.
Library Arrayed by: Greg Lennon, Ph.D.
sequencing by: Washington University Genome Sequencing Center
distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL, send email to:
mage.llnl.gov
imer: -40UP from Gibco
uality sequence stop: 339.
Location/Qualifiers
1..351
/organism="Homo sapiens"
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/db_xref="taxon:9606"
/clone="IMAGE:3441693"
/tissue_type="glioblastoma (pooled)"
/lab_host="DH10B"
/clone_lib="NCI CGAP Brn23"
/note="Organ: brain; Vector: p7T3D-Pac (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTCCATCTGAGTGGAGCGCGCCGATCTTTTCTTTTCTTTTCTTTT
T 3']; double-stranded cDNA was ligated to Eco RI
adaptors (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of the modified p7T3 vector.
Library is normalized, and was constructed by Bento
Soares and M. Fatima Bonaldo."

15.4%; Score 211; DB 10; Length 351;
arity 99.6%; Pred. No. 5.8e-96;
onservative 0; Mismatches 1; Indels 0; Gaps 0;
CCACGCCGACCTCCACTCCTAGCTCCCAATCCCTGACCCCTTGAGGCCCCC 1107
CCACGCCGACCTCCACTCCTAGCTCCCAATCCCTGACCCCTTGAGGCCCCC 292
ATCTCGACTCCCGCTGGCCAGACACCCCGAGGCGATTGTGTCACTGTACTCTGT 1167
ATCTGACTCCCGCTGGCCAGACACCCCGAGGCGATTGTGTCACTGTACTCTGT 232
AAGGATGGTCCAGAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGCGC 1227
AAGGATGGTCCAGAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGCGC 172
AAGCAAGAGACTGGGCTTAGCCAGGAGTCCCAATGTGAGGGGCGAGAACAA 1287
AAGCAAGAGACTGGGCTTAGCCAGGAGTCCCAATGTGAGGGGCGAGAACAA 112
AAGCTCCCTCCCTTGAGAAAT 1309
AAGCTCCCTCCCTTGAGAAAT 90

243 397 bp mRNA linear EST 17-DEC-1999
01.x1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:2315088 3',
sequence.
243

AI69243.1 GI:4834017
EST.
KEYWORDS Homo sapiens (human)
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 397)
REFERENCE
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (Tumor Gene Index)
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Mi
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencin
Clone distribution: NCI-CGAP clone distribution informati
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 438 Std Error: 0.00
Seq primer: -40UP from Gibco.
Location/Qualifiers
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/lab_host="DH10B"
/clone_lib="NCI_CGAP_Pr28"
/note="Organ: prostate; Vector: p7T3D-Pac (Pharm
with a modified polylinker; plasmid DNA from the
normalized library NCI CGAP Pr22 was prepared, an
circles were made in vitro. Following HAP purific
this DNA was used as tracer in a subtractive hybr
reaction. The driver was PCR-amplified cDNAs fro
of 5,000 clones made from the same library (clon
985608-986759, 110192-1101959, and 1217928-1220
Subtraction by Bento Soares and M. Fatima Bonald

Query Match 15.1%; Score 207; DB 9; Length 397;
Best Local Similarity 99.4%; Pred. No. 6.6e-94;
Matches 307; Conservative 0; Mismatches 2; Indels 0;
QY 1065 TCCACCTCAGTCTCCCAATCCCTGACCCCTTGAGGCCCCCAGTGTCTCGAC'
DB 343 TCCACCTCAGTCTCCCAATCCCTGACCCCTTGAGGCCCCCAGTGTCTCGAC'
QY 1125 CCTGGCCACAGACCCCGAGGGCATTGTGTCACTGTACTCTGTGGCAAGGATGG
DB 283 CTGGCCACAGACCCCGAGGGCATTGTGTCACTGTACTCTGTGGCAAGGATGG
QY 1185 GAAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGCGGCGAGGCCAAAG
DB 223 GAAGACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGCGGCGAGGCCAAAA
QY 1245 GGGCCCTAGCCAGGAGTTCCTCCAAATGTGAGGGGCGAGAACCAAGCAAGCTCCTC
DB 163 GGGCCCTAGCCAGGAGTTCCTCCAAATGTGAGGGGCGAGAACCAAGCAAGCTCCTC
QY 1305 AGAATTCCTCTGGGATTTTAAACAGATATTATTTTATTATTATTATTGTGACAAA
DB 103 AGAATTCCTCTGGGATTTTAAACAGATATTATTTTATTATTATTATTGTGACAAA
QY 1365 GATAAATGG 1373
DB 43 GATAAATGG 35

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4 26F1 NTH_MGC_118 Homo sapiens cDNA clone IMAGE:5217367 5',
sequence.
4 4.1 GI:16171193
piens (human)
piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 894)
http://mgs.nci.nih.gov/
1 Institutes of Health, Mammalian Gene Collection (MGC)
shed (1999)
: Robert Strausberg, Ph.D.
cgabbs@mail.nih.gov
Procurement: Life Technologies, Inc.
Library Preparation: Life Technologies, Inc.
Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
Sequencing by: Incyte Genomics, Inc.
Distribution: MGC clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
image.llnl.gov
LLM11546 row: d column: 08
ality sequence start: 5
ality sequence stop: 460.
Location/Qualifiers
1. 894
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5217367"
/tissue_type="leukocyte"
/lab_host="DH10B"
/clone_lib="NIH_MGC_118"
/notes="Vector: pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV
(destroyed); RNA source leukocytes from anonymous pool of
non-activated adult donors. Library is oligo-dT primed
and directionally cloned (EcoRV site is destroyed upon
cloning). Average insert size 1.7 kb, insert size range
1.2-3.3 kb. Library is normalized and enriched for
full-length clones and was constructed by C. Gruber
(Invitrogen). Research Genetics tracking code 027. Note:
this is a NIH_MGC Library."
15.0%; Score 206; DB 12; Length 894;
urity 100.0%; Pred. No. 2.5e-93;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
AGGACGAGACCGCTCGGAACCTGAATCCCGACAGAAAGCCAGGATCTCTGG 351
AGGACGAGACCGCTCGGAACCTGAATCCCGACAGAAAGCCAGGATCTCTGG 281
CTGTAAACGACTAGTTCGCTCGCAGAGTGCACCTAAAGCCGCGAAACACGG 411
CTGTAAACGACTAGTTCGCTCGCAGAGTGCACCTAAAGCCGCGAAACACGG 341
AAGAGCGATCGAGCCCATTTATGAAGTTTCATCCAGCAGCTGACAGGACGAGCG 471
AAGAGCGATCGAGCCCATTTATGAAGTTTCATCCAGCAGCTGACAGGACGAGCG 401
AGGTGTGACGGGACAGTGAG 497
AGGTGTGACGGGACAGTGAG 427
41 5.x1 Soares_NSF_F8_9W_OT_PA_P_S1 Homo sapiens cDNA clone
EST 18-AUG-1998

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IMAGE:1651448 3' similar to contains MSRI.L3 MSRI repetitive
element ;, mRNA sequence.
ACCESSION AI091441
VERSION AI091441.1 GI:3430500
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutelec
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 465)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (C
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs@mail.nih.gov
This clone is available royalty-free through LLNL; contact
IMAGE Consortium (info@image.llnl.gov) for further informat
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 465.
Location/Qualifiers
1. 465
/organism="Homo sapiens"
/mol_type="mRNA"
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/clone="IMAGE:1651448"
/lab_host="DH10B"
/clone_lib="Soares NSF F8_9W_OT_PA_P_S1"
/notes="Organ: pooled; Vector: p773D-Pac (Pharmaci
a modified polylinker; Site 1: Not I; Site 2: Eco
Equal amounts of plasmid DNA from five normalized
libraries were mixed, and ss circles were made in
Following HAP purification, this DNA was used as t
a subtractive hybridization reaction. The driver
PCR-amplified cDNAs from pools of 5,000 clones ma
the same 5 libraries. The pools consisted of the
libraries and clones: Soares NBHSF pool 1:
309384-310919, 323208-325895 Soares NB2HP pool 1:
145032-147335, 147720-148103, 148872-149255, 1500:
150407, 151176-152327 Soares NB2HF8-9W pool 1:
758280-760583, 772104-774407 Soares NBHPA pool 1:
304776-306311, 320136-322823, 326280-326663 Soares
pool 1: 723720-726407, 739080-740999 Subtraction
Soares and M. Fatima Bonaldo."
ORIGIN
Query Match 14.7%; Score 202; DB 9; Length 465;
Best Local Similarity 100.0%; Pred. No. 2.4e-91;
Matches 202; Conservative 0; Mismatches 0; Indels 0; G
QY 25 TCGGCGCGCGGCTCCCGCTCCCGGATCCCTCCCGGATCCCGGATCGGGGGCGGT
Db 183 TCGGCGCGCGGCTCCCGCTCCCGGATCCCTCCCGGATCCCGGATCGGGGGCGGT
QY 85 CAGGCAACGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG
Db 243 CAGGCAACGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG
QY 145 GGGAGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG
Db 303 GGGAGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG
QY 205 TGCTCGGCGCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT
Db 363 TGCTCGGCGCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT
RESULT 62
BF222608/c
LOCUS BF222608
DEFINITION 224 bp mRNA linear EST 09
7p56d12.x1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:364
mRNA sequence.
ACCESSION BF222608

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library Arrayed by: The I.M.A.G.E. Consortium (LINL)
sequencing by: Agencourt Bioscience Corporation
distribution: MGC clone distribution information can be
through the I.M.A.G.E. Consortium/LINL at:
image.linl.gov

row: e column: 12
quality sequence stop: 249.
Location/Qualifiers
1. .1064
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/db_xref="taxon:9606"
/clone="IMAGE:30520331"
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/lab_host="DH10B Tona"
/clone_lib="NIH_MGC_14"
/note="Organ: Placenta; Vector: pBluescriptR; Site:1:
ali-XhoI; Site 2: BamH; Oligo-dr primed using primer
5'-TTTTTTTTTTTTTTTNN-3', size-selected for average
insert size 2.3 kb and normalized to ROT 5. This is a
primary library enriched for full-length clones and
constructed using the Cap-trapper method (Carninci, in
preparation). Library constructed by M. Brownstein
(NIMH/NHGRI, National Institutes of Health). Note: This is
a NIH MGC library."

```

12.7%; Score 175; DB 14; Length 1064;
rity 100.0%; Pred. No. 1.7e-77;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
CCCCCGCCCCATGGCCGCCCTGCGAGCCAGAGCGAGGGCGCCGCGGGGA 149
CCCCCGCCCCATGGCCGCCCTGCGAGCCAGAGCGAGGGCGCCGCGGGGA 87
GGACCGCCCTGCTGGTCCCGCTCGCGCTCGGGCTCGGGCTGGCGCTGCGCT 209
GGACACCGGCCCTGCTGGTCCCGCTCGCGCTGGCGCTGGCGCTGCGCT 147
TTCCTGCTGGCGCGTGGTCAGTTTGGGAGCCGGGCAATCGCTGCCCGAC 264
TCTCTGCTGGCGCGTGGTCAGTTTGGGAGCCGGGCAATCGCTGCCCGAC 202

33 1319 bp mRNA linear EST 26-SEP-2003
 SUBMIT 15623743 NIH MGC 147 Homo sapiens cDNA clone
 00527869 5', mRNA sequence.
 33 33.1 GI:36348525
 sapiens (human)
 sapiens
 Metazoa; Chordata; Vertebrata; Euteleostomi;
 Actinoptera; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
 Accession numbers: 1 to 1319
 Submitter's URL: <http://mgc.hci.nih.gov/>.
 Description: NIH Institutes of Health, Mammalian Gene Collection (MGC)
 Project, released (1999).
 Contact: Daniela S. Gerhard, Ph.D.,
 National Cancer Institute / NIH
 Biotechnology Resource Project, Room 3B10
 Bethesda, MD 20892
 Email: cgapbs-remail.nih.gov
 Procurement: Dr. Stefan Hansson
 Laboratory Preparation: Michael J. Brownstein (NHGRI) with help
 from Piero Carninci (RIKEN)
 Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 Sequencing by: Agencourt Bioscience Corporation
 Distribution: MGC clone distribution information can be
 obtained through the I.M.A.G.E. Consortium/LLNL at:
image.llnl.gov

Plate: NDAM612 row: O column: 14
High quality sequence stop: 287.
Location/Qualifiers
1. 1319
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/db_xref="taxon:9606"
/clone="IMAGE:30527869"
/tissue_type="Human Placenta"
/lab_host="DH10B TonaA"
/clone_lib="NIH MGC 147"
/note="Organ: placenta; Vector: pBluescriptR; Site
all: XhoI; Site 2: BamH; Oligo-dT primed using
5'-TTTTTTTTTTTTTTTN-3', size selected for average
insert size 2.3 kb and normalized to 50. This is
primary library enriched for full-length clones and
constructed using the Cap-trapper method (Carninci
preparation). Library constructed by M. Brownstein
(NIH/NHGRI, National Institutes of Health). Note:
a NIH MGC library."

ORIGIN

Query Match	12.7%;	Score 175;	DB 14;	Length 1319;
Best Local Similarity	100.0%;	Pred. No. 1.8e-77;		
Matches 175;	Conservative 0;	Mismatches 0;	Indels 0;	G+
QY	90	ACAGCCCCCGCCCCCATGTGGCCGCGCGTGGAGCGACAGGCGGAGGGGGCGCGCGG		
Db	29	ACAGCCCCCGCCCCCATGTGGCGCGCCCGTGGAGCGACAGGCGGAGGGGGCGCGCGG		
QY	150	GCGGGGACCGCCCTGTGCTGGTCCCGCTCGCGCTGGGCGCTGGGCGCTGGCGCTGGCGCTT		
Db	89	GCGGGGACCGCCCTGTGCTGGTCCCGCTCGCGCTGGGCGCTGGGCGCTGGCGCTGGCGCTT		
QY	210	CGGCTCTCTCTGGCCGCTGGTTCAGTTTGGGAGCGCGGSCATCCGCTGTCCGCCCCAG ;		
Db	149	CGGCTCTCTCTGGCCGCTGGTTCAGTTTGGGAGCGCGGSCATCGCTGTCCGCCCCAG ;		
RESULT 66				
AA913913/c			338 bp	linear EST 24
LOCUS	AA913913			
DEFINITION	ol35h12.s2 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone			
	IMAGE:1525511.3', mRNA sequence.			

RESULT 66

AA913913/c 338 bp mRNA linear EST 24
LOCUS
DEFINITION
0135h12.s2 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:11525511 3', mRNA sequence.
AA913913
AA913913.1 GI:3053305
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 338)
NCI-CCGAP Project://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project ()
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: ccgaps-rc@mail.nih.gov
This clone is available royalty-free through LLNL ; contac
IMAGE Consortium (info@image.llnl.gov) for further informa
Insert Length: 330 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 332.

FEATURES

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1. 339
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/clone="IMAGE:1525511"
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/clone_lib="soares_NFL_T_GBC_S1"
/notes="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia modified polylinker; Site 1: Not I; Site 2: Eco

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/note="Organ: pancreas; Vector: pBluescript SK-; Site 1: XhoI; Site 2: EcoRI; Constructed with lambda ZAPII system (Stratagene) by Dr. J. Ferrer, in vivo mass-excised to pBluescript SK- by Dr. H. Inoue following the Washington University protocol (http://genome.wustl.edu/est/lambda_protocol.shtml). Please contact Hiroshi Inoue, MD/PhD for further information on this library (Metabolism Division, Permutt Laboratory, Washington University School of Medicine, Box 8127, 660 S Euclid Ave, St. Louis, MO 63110). Note: this is a Washington University Pancreas EST project library."

12.2%; Score 167; DB 12; Length 422;
rity 100.0%; Pred. No. 1.7e-73; Indels 0; Gaps 0;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
TGCCCTGGGCGCTTTCACGTGTTTCCATCCACATAAATACAGTATCCCACTCT 1036
TGCCCTGGGCGCTTTCACGTGTTTCCATCCACATAAATACAGTATCCCACTCT 363
TACAACTCCCGACGGCCACTTCCACCTCACTAGCTCCCAATCCCTGACCCCT 1096
TACAACTCCCGACGGCCACTTCCACCTCACTAGCTCCCAATCCCTGACCCCT 303
GCCCGCAGTGATCTCGACTCCCGCTGGCCACAGACCCCGCAG 1143
GCCCGCAGTGATCTCGACTCCCGCTGGCCACAGACCCCGCAG 256

6 372 bp mRNA linear EST 17-SEP-2001
6 y1 Human insulinoma Homo sapiens cDNA 5', mRNA sequence.

6.1 GI:15630163

apiens (human)

ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Homnidae; Homo.
es 1 to 372)
D., Brown, J., Kent, G., Permutt, A., Lee, C., Kaestner, K.,
ca, I., Seearce, M., Brestelli, J., Gradwohl, G., Clifton, S.,
L., Marra, M., Page, D., Wylie, T., Martin, J., Blistain, A.,
A., Theising, B., Ritter, S., Ronko, I., Bennett, J.,
as, M., Gibbons, M., McCann, R., Cole, R., Tsagarelshvili, R.,
ns, T., Jackson, Y., and Bowers, Y.
ine Pancreas Consortium
ished (2000)
ESTs: id87a02.x1

2: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
ine Pancreas Consortium
a University, Howard Hughes Medical Institute
f Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
38

17-495-1812
17-495-8557
dnelton@biohp.harvard.edu
y was constructed by Dr. J. Ferrer In vivo mass-excised to
cript SK- by Dr. H. Inoue DNA sequencing by: Washington
sity Genome Sequencing Center For information on obtaining a
Please contact: Dr. Hiroshi Inoue (hinoue@im.wustl.edu)
imer: -40RP from Gibco.

Location/Qualifiers
1..372
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XhoI; Site 2: EcoRI; Constructed with lambda ZAPII
(Stratagene) by Dr. J. Ferrer, in vivo mass-excise
pBluescript SK- by Dr. H. Inoue following the Was
University protocol

(http://genome.wustl.edu/est/lambda_protocol.shtml)
Please contact Hiroshi Inoue, MD/PhD for further
information on this library (Metabolism Division,
Laboratory, Washington University School of Medic
8127, 660 S Euclid Ave, St. Louis, MO 63110). Not
is a Washington University Pancreas EST project li

ORIGIN

Query Match 11.8%; Score 162; DB 12; Length 372;
Best Local Similarity 100.0%; Pred. No. 6e-71;
Matches 162; Conservative 0; Mismatches 0; Indels 0; G
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DB 211 GCATTGTGTTCACTGCTCTGTGGCAAGGATGGTCCAGAGACCCCACTTCAG
QY 1205 TAAGAGGGGCTGGACCTGTGGCGCAGGAAGCCAAAGAGACTGGGCGCTAGGCCAGGAG
DB 271 TAAGAGGGGCTGGACCTGTGGCGCAGGAAGCCAAAGAGACTGGGCGCTAGGCCAGGAG
QY 1265 CAAATGTAGGGGGCGAGAAACAGACCAAGCTCTCCCTTGAG 1306
DB 331 CAAATGTAGGGGGCGAGAAACAGACCAAGCTCTCCCTTGAG 372

RESULT 70

AQ890280/c
LOCUS
DEFINITION
HS 3188 B1 F05 MR CIT Approved Human Genomic Sperm Library
sapiens genomic clone Plate=3188 Col=9 Row=L, genomic sur
sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

AQ890280
AQ890280.1 GI:6346470
GSS.
Homo sapiens (human)

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 436)
Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzm
Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams
Hood, L.

TITLE

Sequence-tagged connectors: A sequence approach to mapping
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589
10449764

JOURNAL

MEDLINE
PUBMED
COMMENT
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887

Email: jwallace@u.washington.edu
Clones may be purchased from Research Genetics (info@resge
BAC end Web Server: <http://www.htsc.washington.edu>
Plate: 3188 Row: L column: 9
Seq primer: M13 Reverse
Class: BAC ends
High quality sequence stop: 436.

FEATURES

Location/Qualifiers

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|TGATCTCACTCCCTCCCTGGCCACAGACCCCAAGGCAATTTGTTCACTGTACTCT 141
|GCAAGGATGGTCCAGAGACCCCACTTCAAGGCAATTTGTTCACTGTACTCT 1225
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279 409 bp mRNA linear EST 27-OCT-1999
14.x1 NCI_CGAP_Gas4 Homo sapiens cDNA clone IMAGE:2622510 3',
sequence.
279
279.1 GI:6132886

sapiens (human)
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
lia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
ases 1 to 409)
AP http://www.ncbi.nlm.nih.gov/ncicgap.
nal Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Gene Index
lished (1997)
ct: Robert Strausberg, Ph.D.
: cgapbs-x@mail.nih.gov
a Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
t-Buck, M.D., Ph.D.
Library Preparation: Life Technologies, Inc.
Library Arrayed by: Greg Lennon, Ph.D.
Sequencing by: Washington University Genome Sequencing Center
e distribution: NCI-CGAP clone distribution information can be
through the I.M.A.G.E. Consortium/LLNL at:
io.llnl.gov/bbrp/image/image.html
rimer: -40UP from Gibco
quality sequence stop: 405.
Location/Qualifiers
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ORIGIN

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Matches 158; Conservative 0; Mismatches 0; Indels 0; G

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DB 124 GGCAGAGAAACAGACAGCTCTCCCTTGAGAAATCCCTGTGGATTTTAAACAG
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RESULT 72

R55379 345 bp mRNA linear EST 2;
LOCUS YJ77a08.r1 Soares breast 2NBHst Homo sapiens cDNA clone
DEFINITION IMAGE:154742 5', mRNA sequence.

ACCESSION

R55379

VERSION R55379.1 GI:824674

KEYWORDS EST.

SOURCE Homo sapiens (human).

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 345)

AUTHORS Hillier, L., Clark, N., Dubucque, T., Elliston, K., Hawkins, M.,
Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra,
Parsons, J., Rittkin, L., Rohlfing, T., Soares, M., Tan, F.,
Trevas, E., Waterston, R., Williamson, A., Wohldmann, P., et
Wilson, R.

The WashU-Merck EST Project

Unpublished (1995)

Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

Insert Size: 659

High quality sequence stops: 235 Source: IMAGE Consortium

This clone is available royalty-free through LLNL; conta

IMAGE Consortium (info@image.llnl.gov) for further inform

Insert Length: 659 Std Error: 0.00

Seq primer: M13RP1

High quality sequence stop: 235.

Location/Qualifiers

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/clone="IMAGE:154742"

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/clone_lib="Soares breast 2NBHst"

/note="Organ: breast; Vector: p77T3D (Pharmacia)

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double-stranded cDNA was ligated to Eco RI adapt

(Pharmacia), digested with Not I and cloned into

and Eco RI sites of a modified p77T3 vector (Pha

Library went through one round of normalization

GenCore version 5.1.1.6
copyright (c) 1993 - 2004 Compugen Ltd.

c search, using sw model

il 8, 2004, 19:06:21 ; Search time 654 Seconds
(without alignments)
8918.618 Million cell updates/sec

09-245-198A-3

3
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GO NUC

-op 60.0 , Gapext 60.0

3863 segs, 2124099041 residues

s satisfying chosen parameters: 6747726

lth: 0

lth: 2000000000

sting first 100 summaries

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Geneseqn1980s:*

Geneseqn1990s:*

Geneseqn2000s:*

Geneseqn2001as:*

Geneseqn2001bs:*

Geneseqn2002s:*

Geneseqn2003as:*

Geneseqn2003bs:*

Geneseqn2003cs:*

Geneseqn2004s:*

the number of results predicted by chance to have a
t than or equal to the score of the result being printed,
ed by analysis of the total score distribution.

SUMMARIES

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3.6	1306	7	ACC57901	ACC57901 Human TWE	
3.6	1306	9	ADC35205	ADC35205 Human CDN	
0.8	1364	6	ABK34881	ABK34881 Human CDN	
5.4	1353	3	AAA49717	AAA49717 Human PRO	
5.4	1353	6	ABK40255	ABK40255 cDNA enco	
5.4	1421	2	AAV56000	AAV56000 Human tum	
9.8	1236	2	AAV47613	AAV47613 TNF relat	
9.8	1236	4	AAV47613	AAV47613 Human TRE	
7.1	1030	2	AAV47613	AAV47613 Human TRE	
5.5	898	4	AAV47613	AAV47613 Human TRE	
9.6	493	8	AAV47613	AAV47613 Human TRE	
3.0	195	6	ABK29540	ABK29540 Colon ade	
5.5	282	2	AAV22190	AAV22190 Human gen	
4.7	1239	9	AAV22190	AAV22190 Murine FL	
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27	46	3.4	1168	2	AAV18599	AAV18599 Mus
28	36	2.6	408	7	ABX37032	ABX37032 Bot
29	29	2.1	412	9	ABV56326	ABV56326 Tox
30	26	1.9	26	6	ABK40356	ABK40356 Rev
31	26	1.9	140	7	ABZ78835	ABZ78835 Tun
32	26	1.9	140	7	ABZ09382	ABZ09382 Hun
33	26	1.9	145	7	ABZ78308	ABZ78308 Tun
34	26	1.9	145	7	ABZ78428	ABZ78428 Tun
35	26	1.9	145	7	ABZ08855	ABZ08855 Hun
36	26	1.9	145	7	ABZ08975	ABZ08975 Hun
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40	26	1.9	386	6	ABO60530	ABO60530 Hun
41	26	1.9	391	5	AAH83337	AAH83337 Hun
42	26	1.9	425	7	ABX74646	ABX74646 Hun
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45	26	1.9	452	3	AAV06922	AAV06922 Hur
46	26	1.9	469	6	ABV86720	ABV86720 Hur
47	26	1.9	483	6	ABK55088	ABK55088 Hur
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51	26	1.9	516	3	AAV01272	AAV01272 Hur
52	26	1.9	531	6	ABK55407	ABK55407 Hur
53	26	1.9	540	3	AAV01271	AAV01271 Hur
54	26	1.9	570	3	AAV77898	AAV77898 Hur
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65	22	1.6	263	4	AAV08821	AAV08821 Hu
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77	21	1.5	21	2	AAV18612	AAV18612 SY
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88	20	1.5	20	2	AAV18604	AAV18604 SY
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94	20	1.5	47	3	AAA49772	AAA49772 Hu
95	20	1.5	47	6	ABK40467	ABK40467 In
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AGGATGGGTCCAGAAGACCCCCACTTTCAGGCACCTAAGAGGGGCTGGACCTGGCGGCA 1140
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Standard: cDNA: 1306 BP.

(first entry)

coding sequence.

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/product= "Human TW
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-A2.

2002WO-US023782

2001US-0307838P.

N GENOME SCI INC.

Rosen CA:

0659/40.
2315.

oligomeric complex having a first polypeptide member of the TNF family, a second different member of the TNF family, useful for treating cancer, osteoporosis or an autoimmune disease.

Page 367-368: 388pp: English.

sequence is that of a polynucleotide encoding human TNFAIP3. The invention relates to compositions comprising heterotrimeric complexes of TNF ligand family members, and their use in the treatment of disease, in prevention and treatment of disease. In one embodiment, the heterotrimeric complex comprises full-length or extracellular TNFAIP3 and full-length or extracellular portions of other TNF ligand family members, preferably VEGI or VEGI-SV. The heterotrimeric complex is useful for treating an autoimmune disease, cancer, or other disease, and for inducing apoptosis of cancer cells, increasing B cell proliferation, or inducing apoptosis of cancer cells.

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idard; cDNA; 1364 BP.

(first entry)

ncoding secreted protein #19.

ed protein; gene; ss; nutritional supplement; haemophilia;
 ion; bacterial infection; fungal infection; diabetes; asthma;
 isorder; rheumatoid arthritis; multiple sclerosis; tumour;
 hydritis; allergic reaction; neurodegenerative disease;
 disease; Parkinson's disease; liver fibrosis; cancer; ulcer;
 disorder; inflammatory disorder; Crohn's disease; incision;
 eration; wound healing; burn; haematopoiesis;
 deficiency; lymphoid cell deficiency.

A2.

2001WO-US010224.

2000US-0195582P.

TICS INST INC.

ark HF, Fecthel K, Agostino MJ, Howes SH, Resnick RJ;
Graham JR;

9321/23.

l and ninety two polynucleotides derived from a variety of
 ; sources which encode secreted proteins, useful for treating
 iencies and disorders such as autoimmune disorders.

ie 82; 372pp; English.

n relates to 592 polynucleotides which have been derived from
 ; human tissue sources and which encode novel secreted
 ie polynucleotides can be used as probes for the
 on and isolation of full length cDNA and genomic DNA. The
 des and proteins can also be used as nutritional supplements.

The proteins are useful in the treatment of various immune defici
 and disorders such as viral infections, bacterial infections, fun
 infections, autoimmune disorders (e.g. rheumatoid arthritis, mult
 sclerosis, autoimmune thyroiditis and diabetes) and allergic reac
 and conditions (e.g. asthma). They are also useful for treating
 neurodegenerative diseases (e.g. Alzheimer's disease, Parkinson's
 disease), liver fibrosis, coagulation disorders (e.g. haemophilia
 inflammatory disorders (e.g. Crohn's disease) and tumours. They a
 useful for tissue regeneration, for wound healing and in the trea
 burns, incisions and ulcers. The proteins are also useful for reg
 haematopoiesis and for treating myeloid or lymphoid cell deficien
 Sequences ABK34863-ABK34864 represent polynucleotides of the inve

Sequence 1364 BP; 246 A; 461 C; 394 G; 263 T; 0 U; 0 Other;

Query Match 90.8%; Score 1247; DB 6; Length 1364;
 Best Local Similarity 99.9%; Pred. No. 0;
 Matches 1347; Conservative 0; Mismatches 2; Indels 0; C

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 TTGTGACAAATGTTGATAAATGG 1353

lard; cDNA; 1353 BP.

(first entry)

>DNA clone DNA30879-1152.

; antitumour; tumour; therapy; cytostatic; breast cancer;
 ; renal cancer; colorectal cancer; uterine cancer;
 ar; lung cancer; bladder cancer;
 is system cancer; melanoma; leukaemia; neoplasia; ss.

Location/Qualifiers

58..807

/*tag= a

58..177

/*tag= b

178..804

/*tag= c

2.

99WO-US028565.

98US-0113296P.

99WO-US005028.

99US-0130232P.

28-APR-1999; 99US-0131445P.
 14-MAY-1999; 99US-0134287P.
 20-JUL-1999; 99US-0144758P.
 26-JUL-1999; 99US-0145698P.
 15-SEP-1999; 99WO-US021090.
 15-SEP-1999; 99WO-US021547.
 (GETH) GENENTECH INC.
 Ashkenazi AJ, Goddard A, Godowski PJ, Gurney AL, Marsters SA;
 Napier MA, Pitti RM, Wood WI;
 P-PSDB; AAY95338.
 WPI; 2000-442668/38.
 Novel composition to inhibit neoplastic cell growth or for treatin
 in mammal comprises polypeptides PRO179, PRO207, PRO320, PRO219, F
 PRO224, PRO328, PRO301, PRO526, PRO362, PRO356, PRO509 or PRO866.
 Claim 20; Fig 3; 172pp; English.
 The present sequence is that of cDNA clone DNA30879-1152 (ATCC 209
 encoding human PRO207 (see AAY95338), which shows homology to seve
 members of the tumour necrosis factor family, especially human
 lymphotoxin (23.4%). The cDNA was identified in a foetal kidney cl
 library following identification of an expressed sequence tag wit
 homology to human Apo-2 ligand. A claimed method for inhibiting t
 growth of a tumour cell comprises exposing the tumor cell to PRO1
 PRO207, PRO320, PRO219, PRO224, PRO328, PRO301, PRO526, P
 PRO356, PRO509 or PRO866 (see AAY95337-49), their agonists or chi
 polypeptides incorporating them. The tumour is especially a cancer
 selected from breast, ovarian, renal, colorectal, uterine, prosta
 lung, bladder and central nervous system cancer, melanoma and leu
 Nucleic acids encoding PRO179 etc. are used in the recombinant pr
 of the antitumour polypeptides

Sequence 1353 BP; 257 A; 443 C; 389 G; 264 T; 0 U; 0 Other;

Query Match 85.4%; Score 1172; DB 3; Length 1353;
 Best Local Similarity 99.8%; Pred. No. 0;
 Matches 1322; Conservative 0; Mismatches 3; Indels 0; G

QY	49	CGATCCCTCGGCTCCCGGATGCGGGGGGGGGTGGAGCAGGACAGCCCGCCG
DB	1	CGATCCCTCGGCTCCCGGATGCGGGGGGGGGTGGAGCAGGACAGCCCGCCG
QY	109	GCGGCGGCTCGGAGCCAGAGGCGGAGGCGGCGGCGGAGCGCGGCGGCGGCT
DB	61	GCGGCGGCTCGGAGCCAGAGGCGGAGGCGGCGGCGGAGCGCGGCGGCGGCT
QY	169	GTCGCGCTCGGCTCGGCGCTGGGCGCTGGGCGCTGGGCGCTGGGCGCTGGGCG
DB	121	GTCGCGCTCGGCTCGGCGCTGGGCGCTGGGCGCTGGGCGCTGGGCGCTGGGCG
QY	229	GTCAGTTTGGGAGGCGGCGATCGCTGTCGCGCCGAGGAGCGCTGCCAGGAGGAGCT
DB	181	GTCAGTTTGGGAGGCGGCGATCGCTGTCGCGCCGAGGAGCGCTGCCAGGAGGAGCT
QY	289	GCAGAGGAGGAGCCAGGACCCGTCGGAATCTCCCGAGACAGAGAAAGCCAGGA
DB	241	GCAGAGGAGGAGCCAGGACCCGTCGGAATCTCCCGAGACAGAGAAAGCCAGGA
QY	349	GCGCCTTTCTGAAACCGACTAGTTTCGGCTTCGAGAGTGCACCTAAAGCCGAA
DB	301	GCGCCTTTCTGAAACCGACTAGTTTCGGCTTCGAGAGTGCACCTAAAGCCGAA
QY	409	CGGCTCGAAGAGAGATCGCAGCCCATTTATGAAGTTTATCCACAGCTGGACAGG
DB	361	CGGCTCGAAGAGAGATCGCAGCCCATTTATGAAGTTTATCCACAGCTGGACAGG
QY	469	GCGCAGGACAGGTTGGAGCGGAGCAGTCAAGTGGTGGGAGGAGGAGCAGAAATCAACAG
DB	421	GCGCAGGACAGGTTGGAGCGGAGCAGTCAAGTGGTGGGAGGAGGAGCAGAAATCAACAG

GTTTGGGAGCGGSCATCGCTGTCCGCCAGGAGCCTGCCAGGAGAGCTGGTG 288
 |||||
 GTTTGGGAGCGGSCATCGCTGTCCGCCAGGAGCCTGCCAGGAGAGCTGGTG 274
 |||||
 AGGAGGACACGACCCGTCGGAACCTGAATCCCCAGACAGAAAGACGAGATCCT 348
 |||||
 AGGAGGACACGACCCGTCGGAACCTGAATCCCCAGACAGAAAGACGAGATCCT 334
 |||||
 CTCTTCTGAACCGACTAGTTCGGCTCGCAGAAAGTGACCTAAAGCCGGAACA 408
 |||||
 CTCTTCTGAACCGACTAGTTCGGCTCGCAGAAAGTGACCTAAAGCCGGAACA 394
 |||||
 CTGAGAGAGCGATCCGACCCATTAATGAAGTTTCACACGACCTGACAGACGGA 468
 |||||
 CTGAGAGAGCGATCCGACCCATTAATGAAGTTTCACACGACCTGACAGACGGA 454
 |||||
 AGGCAGGTGTGACCGGACAGTGAAGTGGGAGGAAGCCAGAAATCAACAGTCC 528
 |||||
 AGGCAGGTGTGACCGGACAGTGAAGTGGGAGGAAGCCAGAAATCAACAGTCC 514
 |||||
 CTCTCGGCTACAAACCGCAGATCGGGAGTTTATAGTCACCGGGCTGGCTCTAC 588
 |||||
 CTCTCGGCTACAAACCGCAGATCGGGAGTTTATAGTCACCGGGCTGGCTCTAC 574
 |||||
 CTGTACTGTGACGTCGACTTTGATGAGGGGAAGCTGTCTACCTGAAGCTGGACTTG 648
 |||||
 CTGTACTGTGACGTCGACTTTGATGAGGGGAAGCTGTCTACCTGAAGCTGGACTTG 634
 |||||
 GTGGATGTGTGTGGCCCTCGCTGCTGGAGGAATTCAGCCACTGCGGCGAGT 708
 |||||
 GTGGATGTGTGTGGCCCTCGCTGCTGGAGGAATTCAGCCACTGCGGCGAGT 694
 |||||
 CTCGGGCCCAGCTCCGCTCTGCGAGTGTCTGGGCTGTGGCCCTGCGGCGAGG 768
 |||||
 CTCGGGCCCAGCTCCGCTCTGCGAGTGTCTGGGCTGTGGCCCTGCGGCGAGG 754
 |||||
 TCCCTGGGATCCGACCCCTCCCTGCGGCCATCTCAAGGCTGCGCCCTTCTCACC 828
 |||||
 TCCCTGGGATCCGACCCCTCCCTGCGGCCATCTCAAGGCTGCGCCCTTCTCACC 814
 |||||
 TTCCGACTCTTCCAGGTTCACTAGGAGGCGCTGTCTCCCAAGTGTCTCCAGGCT 888
 |||||
 TTCCGACTCTTCCAGGTTCACTAGGAGGCGCTGTCTCCCAAGTGTCTCCAGGCT 874
 |||||
 GGCTCCCTCGACAGCTCTCTGGGACCCGCTCCCTCTGCGCCAGCTCTAGGCGCT 948
 |||||
 GGCTCCCTCGACAGCTCTCTGGGACCCGCTCCCTCTGCGCCAGCTCTAGGCGCT 934
 |||||
 TGCTCCAGACCTGCCCCCTCTAGAGGCTGCTGCGGCTGTTCACGTTTTCCTCA 1008
 |||||
 TGCTCCAGACCTGCCCCCTCTAGAGGCTGCTGCGGCTGTTCACGTTTTCCTCA 994
 |||||
 ACATATAATACAGTATCCCACTCTTAATCAACTCCGCCAGCCGCTCTCCA 1068
 |||||
 ACATATAATACAGTATCCCACTCTTAATCAACTCCGCCAGCCGCTCTCCA 1054
 |||||
 TCACTAGTCCCAATCCCTGACCTTTGAGGCCCCCAGTGTCTGACTTCCCTCCCTG 1128
 |||||
 TCACTAGTCCCAATCCCTGACCTTTGAGGCCCCCAGTGTCTGACTTCCCTCCCTG 1114
 |||||
 TACAGACCCCGAGGCTGTTGTTCACTGTCTGTGGCAAGGAGTGGTCCAGAG 1189
 |||||
 TACAGACCCCGAGGCTGTTGTTCACTGTCTGTGGCAAGGAGTGGTCCAGAG 1174
 |||||
 CCCACTTCAGGCACTAAGAGGGGCTGACCTGCGGCGAGAGCCAAAGAGACTGGC 1248
 |||||
 CCCACTTCAGGCACTAAGAGGGGCTGACCTGCGGCGAGAGCCAAAGAGACTGGC 1234
 |||||
 AGGCCAGAGTTCCTCAATGTGAGGCGGAGAACAGACAGCTCTCTCTTGGAGAA 1308
 |||||
 AGGCCAGAGTTCCTCAATGTGAGGCGGAGAACAGACAGCTCTCTCTTGGAGAA 1294
 |||||
 CCCTGTGATTTTAAACAGATATTATTTTATTATTATTGTGACAAATGTTGATA 1368

Db 1295 TTCCCTGTGATTTTAAACAGATATTATTTTATTATTATTCGACAAATGTT
 Qy 1369 AATGG 1373
 Db 1355 AATGG 1359
 |||||
 RESULT 9
 AAV47613
 ID AAV47613 standard; cDNA; 1236 BP.
 XX
 AC AAV47613;
 XX
 DT 27-OCT-1998 (first entry)
 XX
 DE TNF related endothelium proliferative agent gene.
 XX
 KW ss; TNF; endothelium proliferative agent; TREPA; wound healing; tissue grafting; vascularisation; apoptosis; autoimmune; birth of
 XX Homo sapiens.
 XX
 FH Key Location/Qualifiers
 CDS 1..750
 FT /*tag= a
 FT /product= "TREPA"
 FT
 XX WO9835061-A2.
 XX
 PD 13-AUG-1998.
 XX
 PF 12-FEB-1998; 98WO-US002859.
 XX
 PR 12-FEB-1997; 97US-00798692.
 PR 10-FEB-1998; 98US-00021706.
 XX
 PA (ABBO) ABBOTT LAB.
 XX
 PI Wiley SR;
 XX
 DR WPI; 1998-447255/38.
 DR P-PSDB; AAW29745.
 XX
 PT Detecting nucleic acid encoding TREPA - useful for diagnosis and treatment of autoimmune disease, tumours and inflammation.
 PS
 PS Claim 11; Page 123-4; 142pp; English.
 XX
 CC The TNF-related endothelium proliferative agent (TREPA), or its activators or agonists, are used to treat a deficit of TREPA, e promote wound healing or tissue grafting, by promoting vascular also to induce apoptosis for treating cancer and eliminating au T cells, as an adjunct to cancer chemotherapy or antiviral treat TREPA peptides can also be used to target cytotoxic agents or f affinity isolation of the corresponding receptor, the nucleic a which can be used to transform tumour cells to render them more responsive to TREPA and to screen for TREPA mimics. Ribozymes, RNA , antibodies or peptides, are used to treat TREPA-associated diseases, e.g. tumours and metastases (by inhibiting vascularis inflammation or a wide range of autoimmune conditions, conditio involving abnormal stimulation of epithelial cells (e.g. atherosclerosis), for birth control (inhibiting ovulation and p formation) or other angiogenic conditions (e.g. ulcers)
 XX
 SQ Sequence 1236 BP; 225 A; 416 C; 358 G; 237 T; 0 U; 0 Other;
 Query Match 69.8%; Score 958; DB 2; Length 1236;
 Best Local Similarity 99.6%; Pred. No. 0;
 Matches 1208; Conservative 0; Mismatches 5; Indels 0;

Qy 129 GCGAGGGGGCGCCCGGGGAGCCCGGACCCGCTGTGTGTCGCGCTCGCGCT

BP; 223 A; 317 C; 279 G; 211 T; 0 U; 0 Other;
 57.1%; Score 784; DB 2; Length 1030;
 99.9%; Pred. No. 0;
 0; Mismatches 1; Indels 0; Gaps 0;
 TTTGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTG 288
 TTTGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTG 60
 GGAGACACGAGACCGCTCGGAATGAATCCCGAGACAGAGAAAGCCAGGATCCT 348
 GGAGACACGAGACCGCTCGGAATGAATCCCGAGACAGAGAAAGCCAGGATCCT 120
 TTTCTGTAACCGATAGTAGTTCGGCTCGGAGAGTGCATTAAGCCGCGAAACA 408
 TTTCTGTAACCGATAGTAGTTCGGCTCGGAGAGTGCATTAAGCCGCGAAACA 180
 TCGAGAGCGATCGAGCGCCATTATGAATTCATCCAGCTGACAGGACGGA 468
 TCGAGAGCGATCGAGCGCCATTATGAATTCATCCAGCTGACAGGACGGA 240
 AGCAGGTGTGGACGGGACAGTGTGGTGGGAGGAGCCAGAAATCAAGCTCC 528
 AGCAGGTGTGGACGGGACAGTGTGGTGGGAGGAGCCAGAAATCAAGCTCC 300
 TCTGCGCTACAAACCGCAGATCGGAGTGTATAGTACCCGGGCTGGCTCTAC 588
 TCTGCGCTACTACCGCAGATCGGAGTGTATAGTACCCGGGCTGGCTCTAC 360
 TGTACTGTCAAGTGCATTTGATGAGGGAGGCTGTCTACCTGAAGCTGGACTTG 648
 TGTACTGTCAAGTGCATTTGATGAGGGAGGCTGTCTACCTGAAGCTGGACTTG 420
 TGGATGTGTGCTGGCCCTCGCTGCTGAGGAAATTTCTCAGCCACTGCGGCCAGT 708
 TGGATGTGTGCTGGCCCTCGCTGCTGAGGAAATTTCTCAGCCACTGCGGCCAGT 480
 TCGGCGCCAGCTCGGCTCTGCGAGTGTCTGGCTGTGGCTGTGGCCAGG 768
 TCGGCGCCAGCTCGGCTCTGCGAGTGTCTGGCTGTGGCTGTGGCCAGG 540
 TCGTGGGATCCGACCCCTCGCTGGCCATCTCAAGGCTGCGCCCTTCTCTACC 828
 TCGTGGGATCCGACCCCTCGCTGGCCATCTCAAGGCTGCGCCCTTCTCTACC 600
 TCGGACTCTTCCAGGTTCACTGAGGGCCCTGCTCTCCACAGTGTCCAGGCT 888
 TCGGACTCTTCCAGGTTCACTGAGGGCCCTGCTCTCCACAGTGTCCAGGCT 660
 GCTCCCTCGACAGCTCTGTGGGACCGGTCCTCTGCGCCACCTCAGCGCT 948
 GCTCCCTCGACAGCTCTGTGGGACCGGTCCTCTGCGCCACCTCAGCGCT 720
 GCTCCAGCTGCGCCCTCTAGAGGCTGCTGGGCTGTTCACGTGTTTCCA 1008
 GCTCCAGCTGCGCCCTCTAGAGGCTGCTGGGCTGTTCACGTGTTTCCA 780
 ACATAAATACAGTATCCCACTTATCTATACAACTCCCGCCAGCCCACT 1063
 ACATAAATACAGTATCCCACTTATCTATACAACTCCCGCCAGCCCACT 835
 dard; DNA; 898 BP.
 (first entry)
 -ctor pDC409-LZ-TWEAK fusion protein-encoding DNA.

XX TWEAK extracellular domain; tumour necrosis factor; TNF; angiogen
 KW ocular neovascularisation; diabetic retinopathy; neovascular glau
 KW retinoblastoma; retinopathy of prematurity; retrolental fibroplas
 KW rubeosis; uveitis; macular degeneration; arthritis; rheumatism; d
 KW corneal graft neovascularisation; psoriasis; metastatic condition
 KW malignant tumour; sarcoma; carcinoma; benign tumour; haemophilic
 KW preneoplastic condition; myocardial angiogenesis; wound granulat
 KW scleroderma; vascular adhesion; telangiectasia; ischaemia; human
 KW atherosclerotic plaque neovascularisation; coronary atheroscler
 KW peripheral atherosclerosis; pDC409-LZ-TWEAK; TWEAK receptor; TWEA
 fusion protein.
 XX Homo sapiens.
 OS Synthetic.
 XX Location/Qualifiers
 FT CDS 52..873
 FT /tag= a
 FT /product= "Fusion protein comprising a growth hor
 leader, a leucine zipper multimerisation domain,
 human TWEAK extracellular domain"
 XX WO200145730-A2.
 XX 28-JUN-2001.
 XX 19-DEC-2000; 2000WO-US034755.
 XX 20-DEC-1999; 99US-0172878P.
 XX 10-MAY-2000; 2000US-0203347P.
 XX (IMMUNEX CORP.
 XX WILEY SR;
 XX WPI; 2001-417975/44.
 XX P-PSDB; AAU03499.
 XX Modulating angiogenesis in a mammal for treating diseases mediate
 angiogenesis, e.g. solid tumors and vascular deficiencies of card
 peripheral tissue, by administering antagonist or agonist of TWEA
 receptor.
 XX Example 1; Page 39-40; 46pp; English.
 XX The sequence represents a DNA from the expression vector pDC409-L
 which encodes a fusion protein comprising a growth hormone leader
 leucine zipper multimerisation domain, and the extracellular doma
 human TWEAK. The fusion protein was used in the isolation of huma
 receptor (TWEAKR)-expressing clones from a COS cell human cDNA li
 The TWEAK protein is a member of the tumour necrosis factor (TNF)
 and induces angiogenesis. TWEAKR may therefore be used to screen
 develop TWEAKR agonists and antagonists for the modulation of
 angiogenesis, to be used in the treatment and diagnosis of human
 The disorders mediated by angiogenesis include ocular disorders
 characterised by ocular neovascularisation such as diabetic retin
 neovascular glaucoma, retinoblastoma, retinopathy of prematurity,
 retrolental fibroplasia, rubeosis, uveitis, macular degeneration
 corneal graft neovascularisation, and inflammatory diseases such
 arthritis, rheumatism and psoriasis. Other treatable diseases inc
 malignant and metastatic conditions such as sarcomas and carcinom
 benign tumours and preneoplastic conditions, myocardial angiogene
 haemophilic joints, scleroderma, vascular adhesions, atheroscler
 plaque neovascularisation, telangiectasia, wound granulation, cor
 atherosclerosis, peripheral atherosclerosis and ischaemia
 XX Sequence 898 BP; 187 A; 266 C; 267 G; 178 T; 0 U; 0 Other;
 XX Query Match 45.5%; Score 625; DB 4; Length 898;
 Best Local Similarity 100.0%; Pred. No. 6e-281;
 Matches 625; Conservative 0; Mismatches 0; Indels 0; C

TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 291
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 309
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 351
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 369
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 411
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 429
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 471
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 489
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 531
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 549
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 591
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 609
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 651
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 669
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 711
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 729
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 771
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 789
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 831
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 849
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 856
 TGGGAGCGGCGATCGCTGTCCGCCAGGAGCTGCCAGGAGGAGCTGGTGCA 874

Standard; cDNA; 493 BP.

(first entry)

Human cell cDNA #2146.

Sequencing by hybridisation; SBH; expressed sequence tag; EST;
 tag; biodiversity; genetic disorder.

-A1.

2001US-00918995.

2001US-00918995.

NAC R. T.

T. I.

HE-CRAIN B.

SON M C.

S L W.

XX Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;
 XX WPI; 2003-615964/58.
 XX New polynucleotide sequences obtained from various cDNA libraries
 PT as hybridization probes, as oligomers for PCR, for chromosome and
 PT mapping, in the recombinant production of protein, or in generati
 PT antisense DNA or RNA.
 XX Claim 1; SEQ ID NO 21225; 44pp; English.
 XX The invention relates to an isolated polynucleotide comprising an
 CC 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequ
 CC determined by the technique of SBH (sequencing by hybridisation).
 CC included is a purified polypeptide comprising a sequence correspo
 CC a reading frame of the novel polynucleotide. The nucleic acid seq
 CC are useful in diagnostics as expressed sequence tags (EST) for
 CC identifying expressed genes or for physical mapping of the human
 CC in forensics, in assessing biodiversity, or in identifying muta
 CC responsible for genetic disorders and other traits. The nucleotide
 CC sequences are also useful as hybridisation probes, as oligomers f
 CC for chromosome and gene mapping, in the recombinant production of
 CC protein, or in generating antisense DNA or RNA. The purified poly
 CC is useful for generating antibodies specific for it. The present
 CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequen
 CC for this patent did not form part of the printed specification, b
 CC obtained in electronic format directly from USPTO at
 CC seqdata.uspto.gov/sequence.html?docID=20030073623
 XX Sequence 493 BP; 87 A; 180 C; 120 G; 101 T; 0 U; 5 Other;

Query Match 29.6%; Score 407; DB 8; Length 493;
 Best Local Similarity 100.0%; Pred. No. 2.6e-179;
 Matches 407; Conservative 0; Mismatches 0; Indels 0; C

QY 874 CAGTCGTCCAGGCTGCGGCTCCCTCGACAGCTCTCTGGGACCCGGTCCCTC
 DB 87 CAGTCGTCCAGGCTGCGGCTCCCTCGACAGCTCTCTGGGACCCGGTCCCTC
 QY 934 CCACCTCAGCGCTCTTTGCTCCAGACTGCTCCCTCTAGAGGCTGCTG
 DB 147 CCACCTCAGCGCTCTTTGCTCCAGACTGCTCCCTCTAGAGGCTGCTG
 QY 994 TTCAGTGTCTTCCATCCCATATAATACAGTATTCCTTATCTTACAACT
 DB 207 TTCAGTGTCTTCCATCCCATATAATACAGTATTCCTTATCTTACAACT
 QY 1054 ACCGCCACTCTCCACTCAGTCTCCCAATCCCTGACCTTTGAGGCCCCCA
 DB 267 ACCGCCACTCTCCACTCAGTCTCCCAATCCCTGACCTTTGAGGCCCCCA
 QY 1114 CTCGACTCCCCCTGGCCACAGACCCCCCAGGGCAATGTGTTCACTGTACTCTGT
 DB 327 CTCGACTCCCCCTGGCCACAGACCCCCCAGGGCAATGTGTTCACTGTACTCTGT
 QY 1174 GGATGGGTCCAGAGACCCCACTTCAGGCACTAAGAGGGGCTGGACCTGGGGCA
 DB 387 GGATGGGTCCAGAGACCCCACTTCAGGCACTAAGAGGGGCTGGACCTGGGGCA
 QY 1234 CCAAGAGACTGGGCTAGGCGCAGGAGTCCCAATGTGAGGGCGCA 1280
 DB 447 CCAAGAGACTGGGCTAGGCGCAGGAGTCCCAATGTGAGGGCGCA 493

RESULT 14

ABK29540

ID ABK29540 standard; cDNA; 195 BP.

XX AC

XX ABK29540;

XX 23-APR-2002 (first entry)

XX

EAK; TNF relatedness and weak ability to induce cell death;
ecrosis factor; TWEAK; fibrosis; kidney disease;
lung disease; kidney disease; skin disease;
le disease; adipose tissue disease;
nal tract disease; pancreatic disease;
organ disease; neural disease; cartilage disease;
connective tissue disease; cellular death; hepatotropic;
1; gastrointestinal; osteopathic; gene; ss.

Location/Qualifiers
1. .750
/*tag= a
/product= "FL-TWEAK"

A2.

2003WO-US011350.

2002US-0371611P.

IN INC.

kubowski A, Zheng T, Hahn K;

1256/78.

7/12.

WEAK-related condition, e.g. liver, gastrointestinal, kidney,
atic, cartilage or neural tissue condition in a subject
ministering to the subject a TWEAK agonist or antagonist.

3Q ID NO 2; 120pp; English.

sequence is the coding sequence for murine transmembrane FL-
latedness and weak ability to induce cell death, where TNF
rosis factor). TWEAK is a member of the TNF family. TWEAK
antagonists are useful for treating a TWEAK-related
g. fibrosis; cardiac disease; liver disease; lung disease;
se; skin disease; skeletal muscle disease; adipose tissue
ointestinal tract disease; pancreatic disease; reproductive
; neural disease; cartilage disease; bone disease;
issue disease; cellular death; and a pathological condition
expressing a TWEAK receptor.

9 BP; 249 A; 386 C; 331 G; 273 T; 0 U; 0 Other;

4.7%; Score 64; DB 9; Length 1239;

larity 100.0%; Pred. No. 2e-19;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CGCCCGCGTGGAGCCGAGCGGAGGGGGCGCGGGGAGCGGGGACCGGCGCTG 165

CGCCCGCGTGGAGCCGAGCGGAGGGGGCGCGGGGAGCGGGGACCGGCGCTG 60

G 169

G 64

ndard; DNA; 60 BP.

(first entry)

d transcript detection oligonucleotide SEQ ID NO:31596.

Human; mouse; rat; splice transcript; detection; RNA transcript;
splice variant; transcriptome; oligonucleotide library; ss.

Homo sapiens.

WO200210449-A2.

07-FEB-2002.

20-JUL-2001; 2001WO-IB001903.

28-JUL-2000; 2000US-0221607P.

02-MAY-2001; 2001US-0287724P.

(COMP-) COMPUGEN INC.

Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

WPI; 2002-257383/30.

New oligonucleotide libraries comprising oligonucleotides which
selectively hybridize to mRNAs transcribed from a transcription
genome, useful for detecting tissue-, pathology-, and development
specific genes.

Example 1; SEQ ID NO 31596; 47pp; English.

The present invention describes oligonucleotide libraries for det-
messenger RNAs that populate a (sub-)transcriptome, where the (s
)transcriptome comprises messenger RNAs transcribed from multiple
transcription units that populate a genome. The library comprises
oligonucleotides, each capable of hybridising selectively to a s
messenger RNAs transcribed from a given transcription unit of the
which encodes one or more messenger RNA splice variants. The
oligonucleotide libraries are useful for detecting mRNAs from a
biological sample, in expression profiling studies, in qualitati-
quantitatively characterising the corresponding transcriptome, a
detecting RNA transcripts and splice variants of human or animal
transcriptomes. The libraries may also be used as specialised mi-
libraries to detect transcripts of a sub-transcriptome under a p
biological or pathological state, and so allowing the detection
- and pathology-specific genes such as those genes only expresse
specific tissue under a specific pathological condition; to dete
developmental specific genes; and to detect RNA transcripts and
variants of a transcriptome of a patient suffering from a partic
disorder. ABN27253 to ABN59589 represent oligonucleotide sequen-
rats, humans and mice, which are used in the exemplification of
present invention. N.B. The sequence data for this patent did no
part of the printed specification, but was obtained in electroni-
directly from WIPO at ftp.wipo.int/pub/published_pct_sequences

Sequence 60 BP; 13 A; 16 C; 17 G; 14 T; 0 U; 0 Other;

Query Match 4.4%; Score 60; DB 6; Length 60;

Best Local Similarity 100.0%; Pred. No. 1.7e-17;

Matches 60; Conservative 0; Mismatches 0; Indels 0;

1145 GCATTGTGTTCACTGTACTCTGTGGCAAGGATGGGTCCAGAACCCCACTTCA

1 GCATTGTGTTCACTGTACTCTGTGGCAAGGATGGGTCCAGAACCCCACTTCA

RESULT 18

ABN41049

ID ABN41049 standard; DNA; 60 BP.

XX AC ABN41049;

XX 15-JUL-2002 (first entry)

Human spliced transcript detection oligonucleotide SEQ ID NO:137

Human; mouse; rat; splice transcript; detection; RNA transcript;

; transcriptome; oligonucleotide library; ss.

001WO-IB001903.

000US-0221607P.

001US-0287724P.

EN INC.

sserman A, Mintz E, Mintz L, Faigler S;

183/30.

otide libraries comprising oligonucleotides which
hybridize to mRNAs transcribed from a transcription unit of a
for detecting tissue-, pathology-, and developmental-

; ID NO 13797; 47pp; English.

vention describes oligonucleotide libraries for detecting
that populate a (sub-)transcriptome, where the (sub-
comprises messenger RNAs transcribed from multiple
units that populate a genome. The library comprises several
les, each capable of hybridizing selectively to a set of
transcribed from a given transcription unit of the genome,
one or more messenger RNA splice variants. The
le libraries are useful for detecting mRNAs from a
ple, in expression profiling studies, in qualitatively or
transcripts and splice variants of human or animal
; The libraries may also be used as specialised mini
lect transcripts of a sub-transcriptome under a particular
pathological state, and so allowing the detection of tissue
ly-specific genes such as those genes only expressed in
under a specific pathological condition; to detect
specific genes; and to detect RNA transcripts and splice
transcriptome of a patient suffering from a particular
7253 to ABN59589 represent oligonucleotide sequences from
and mice, which are used in the exemplification of the
ion. N.B. the sequence data for this patent did not form
inted specification, but was obtained in electronic format
WIPO at ftp.wipo.int/pub/published_pct_sequences

; 17 A; 15 C; 15 G; 13 T; 0 U; 0 Other;

4.4%; Score 60; DB 6; Length 60;
urity 100.0%; Pred. No. 1.7e-17;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
ATGTGAGGGCGAGAAACAAGACAGCTCCCTTGAGAAATCCCTGTGGATTT 1322
ATGTGAGGGCGAGAAACAAGACAGCTCCCTTGAGAAATCCCTGTGGATTT 60

iard; DNA; 60 BP.

(first entry)

transcript detection oligonucleotide SEQ ID NO:3141.

rat; splice transcript; detection; RNA transcript;
; transcriptome; oligonucleotide library; ss.

XX

OS Homo sapiens.

XX WO200210449-A2.

PN

XX 07-FEB-2002.

PD

XX 20-JUL-2001; 2001WO-IB001903.

PF

XX 28-JUL-2000; 2000US-0221607P.

PR

XX 02-MAY-2001; 2001US-0287724P.

XX (COMP-) COMPUGEN INC.

XX

XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX WPI; 2002-257383/30.

XX

New oligonucleotide libraries comprising oligonucleotides which
selectively hybridize to mRNAs transcribed from a transcription u
genome, useful for detecting tissue-, pathology-, and development
specific genes.

Example 1; SEQ ID NO 31341; 47pp; English.

The present invention describes oligonucleotide libraries for det
messenger RNAs that populate a (sub-)transcriptome, where the (su
)transcriptome comprises messenger RNAs transcribed from multiple
transcription units that populate a genome. The library comprises
oligonucleotides, each capable of hybridising selectively to a se
messenger RNAs transcribed from a given transcription unit of the
which encodes one or more messenger RNA splice variants. The
oligonucleotide libraries are useful for detecting mRNAs from a
biological sample, in expression profiling studies, in qualitativ
quantitatively characterising the corresponding transcriptome, an
detecting RNA transcripts and splice variants of human or animal
transcriptomes. The libraries may also be used as specialised min
libraries to detect transcripts of a sub-transcriptome under a pa
biological or pathological state, and so allowing the detection o
- and pathology-specific genes such as those genes only expressed
specific tissue under a specific pathological condition; to detect
developmental specific genes; and to detect RNA transcripts and s
variants of a transcriptome of a patient suffering from a particu
disorder. ABN27253 to ABN59589 represent oligonucleotide sequenc
rats, humans and mice, which are used in the exemplification of t
present invention. N.B. The sequence data for this patent did not
part of the printed specification, but was obtained in electronic
directly from WIPO at ftp.wipo.int/pub/published_pct_sequences

Sequence 60 BP; 12 A; 19 C; 10 G; 19 T; 0 U; 0 Other;

Query Match 4.4%; Score 60; DB 6; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.7e-17;
Matches 60; Conservative 0; Mismatches 0; Indels 0; G.

QY 979 GGCTGCTGGGCTGTTCACGTGTTTTCATCCACATAAATACAGTATTCACACT
Db 1 GGCTGCTGGGCTGTTCACGTGTTTTCATCCACATAAATACAGTATTCACACT

RESULT 20

ABN58591

ID ABN58591 standard; DNA; 60 BP.

XX

AC ABN58591;

XX

DT 15-JUL-2002 (first entry)

XX

DE Human spliced transcript detection oligonucleotide SEQ ID NO:3133

XX

KW Human; mouse; rat; splice transcript; detection; RNA transcript;

KW splice variant; transcriptome; oligonucleotide library; ss.

XX

98WO-US021407.
97US-0062037P.
97US-0069862P.
TECH INC.
Marsters SA, Pitti R;
982/24.
3- ligand (a tumor necrosis factor) homologue.
je 36; 74pp; English.
vention describes a human tumor necrosis factor (TNF) and
mologue designated Apo-3 ligand. Apo-3 ligand has
ivity. Apo-3 ligand can be used to induce apoptosis in
er cells, to induce NF-kappaB-dependent transcription and
/SAPK-dependent responses in mammalian cells. The present
asents an Apo-3 ligand probe, which is used in an exam
ent invention
P; 10 A; 18 C; 13 G; 9 T; 0 U; 0 Other;
3.6%; Score 50; DB 2; Length 50;
arity 100.0%; Pred. No. 7.7e-13;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
CCCTCTCGCTACACCGCCAGATCGGGGAGTTTATAGTCACCCGG 576
CCCTCTCGCTACACCGCCAGATCGGGGAGTTTATAGTCACCCGG 50
dard; DNA; 50 BP.
(first entry)
DNA probe.
; antitumor; tumor; therapy; cytostatic; breast cancer;
r; renal cancer; colorectal cancer; uterine cancer;
er; lung cancer; bladder cancer;
us system cancer; melanoma; leukaemia; neoplasm; probe; ss.

PA (GETH) GENENTECH INC.
XX Ashkenazi AJ, Goddard A, Godowski PJ, Gurney AL, Marsters SA;
PI Napier MA, Pitti RM, Wood WI;
XX WPI; 2000-442668/38.
XX Novel composition to inhibit neoplastic cell growth or for treati
PT in mammal comprises polypeptides PRO179, PRO207, PRO320, PRO219,
PT PRO224, PRO328, PRO301, PRO526, PRO362, PRO356, PRO509 or PRO866.
XX Example 3; Page 98; 172pp; English.
XX The present sequence is that of a DNA probe based on an isolated
CC expressed sequence tag showing homology to human Apo-2 ligand. Th
CC was used to screen a human foetal kidney cDNA library to identify
CC clone DNA30879-1152 (see AAA49717), which encodes human antitumou
CC protein PRO207 (see AAY95338). A claimed method for inhibiting th
CC of a tumour cell comprises exposing the tumor cell to PRO179, PRO
CC PRO320, PRO219, PRO224, PRO328, PRO326, PRO362, PRO320, P
CC PRO509 or PRO866 (see AAY95337-49). The tumour is especially brea
CC ovarian, renal, colorectal, uterine, prostate, lung, bladder or c
CC nervous system cancer, melanoma or leukaemia. Nucleic acids encod
CC PRO179 etc. are used in the recombinant production of antitumour
CC polypeptides
XX Sequence 50 BP; 10 A; 18 C; 13 G; 9 T; 0 U; 0 Other;
SQ Query Match 3.6%; Score 50; DB 3; Length 50;
Best Local Similarity 100.0%; Pred. No. 7.7e-13; Indels 0; G
Matches 50; Conservative 0; Mismatches 0;
QY 527 CCAGCCCTCTCGCTACACCGCCAGATCGGGGAGTTTATAGTCACCCGG 576
DB 1 CCAGCCCTCTCGCTACACCGCCAGATCGGGGAGTTTATAGTCACCCGG 50
RESULT 24
ABK40292
ID ABK40292 standard; DNA; 50 BP.
XX AC ABK40292;
XX DT 15-JUL-2002 (first entry)
XX DE Oligonucleotide probe for human PRO207 DNA.
XX Human; PRO; benign tumour; malignant tumour; lymphoid malignancy;
KW leukaemia; neuronal disorder; stromal disorder; blastocoele disc
KW inflammatory disorder; immune disorder; angiogenic disorder; cyt
KW neuroprotective; probe; ss.
XX OS Homo sapiens.
XX WO200153486-A1.
XX PD 26-JUL-2001.
XX PF 11-FEB-2000; 2000WO-US003565.
XX PR 08-MAR-1999; 99WO-US005028.
PR 11-MAR-1999; 99US-0123972P.
PR 11-MAY-1999; 99US-0133459P.
PR 02-JUN-1999; 99WO-US012252.
PR 22-JUN-1999; 99US-0140650P.
PR 22-JUN-1999; 99US-0140653P.
PR 20-JUL-1999; 99US-0144758P.
PR 26-JUL-1999; 99US-0145698P.
PR 28-JUL-1999; 99US-0146222P.
PR 17-AUG-1999; 99US-0149395P.
PR 31-AUG-1999; 99US-0151689P.
PR 01-SEP-1999; 99WO-US020111.
PR 15-SEP-1999; 99WO-US021090.

99WO-US028313.
99WO-US028301.
99WO-US028634.
2000WO-US000219.

TECH INC.

Goddard A, Godowski PJ, Gurney AL, Hillan KJ;
Pan J, Pitti RM, Roy MA, Smith V, Stone DM;
Wood WI;

567/26.

nucleic acids encoding PRO polypeptides, useful for treating
ignant tumors, leukemias and lymphoid malignancies,
angiogenic and immunologic disorders.

age 109; 302pp; English.

vention relates to the isolation of novel human PRO

(AAU86128-AAU86162) and the polynucleotide sequences

1. The PRO polypeptides, agonists, antagonists or anti-PRO

useful for treating benign or malignant tumors (e.g.

, bladder, breast, etc), leukemias and lymphoid

other disorders such as neuronal, glial, astrocytal,

glandular, macrophagal, stromal and blastocoeleic disorders,

immune and angiogenic disorders. The polynucleotide

also useful in gene therapy. The present sequence

probe used in the methods of the present invention

1P; 10 A; 18 C; 13 G; 9 T; 0 U; 0 Other;

3.6%; Score 50; DB 6; Length 50;

arity 100.0%; Pred. No. 7.7e-13; Indels 0; Gaps 0;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CCCTCTGGGCTACACCGCCAGATCGGGAGTTTATAGTCACCCGG 576

CCCTCTGGGCTACACCGCCAGATCGGGAGTTTATAGTCACCCGG 50

andard; DNA; 50 BP.

(first entry)

te gene expression profiling probe SEQ ID NO 2558.

3; gene expression profiling; allograft rejection;
sis; congestive heart failure; systemic lupus erythematosus;
thrititis; osteoarthritis; cytomegalovirus; infection; probe;

A2.

2001WO-US047856.

2000US-0241994P.

2001US-0296764P.

ARDIA INC.

, Fry K, Matcuk G, Altman P, Prentice J, Phillips J;
ard R, Quertemous T, Johnson F;

5525/68.

XX
PT
PT
PT
XX
XX
XX

New system for leukocyte expression profiling, diagnosing a disease
monitoring (the rate of) progression of a disease, e.g. atheroscl
or congestive heart failure, comprises diagnostic oligonucleotide
Claim 1; Page 408; Opp; English.

The invention relates to a system for detecting gene expression,
comprises one or two isolated DNA molecules that detect expressio
gene, where the gene corresponds to any of 8143 oligonucleotides
(ABZ00010-ABZ08152) each having 50 base pairs (bp). The system is
for leukocyte expression profiling. It is particularly useful for
diagnosing a disease, monitoring (rate of) progression of a disea
predicting therapeutic outcome, determining prognosis for a patie
predicting disease complications in an individual or monitoring
to treatment in an individual. The diseases include cardiac allo
rejection, kidney allograft rejection, liver allograft rejection,
atherosclerosis, congestive heart failure, systemic lupus erythem
rheumatoid arthritis, osteoarthritis or cytomegalovirus infectio
Sequence 50 BP; 14 A; 11 C; 19 G; 6 T; 0 U; 0 Other;

Query Match 3.6%; Score 50; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 7.7e-13;
Matches 50; Conservative 0; Mismatches 0; Indels 0; C

QY 1196 TTCAGGCACTAAGAGGGGCTGGACCTGGCGGCGAGGAGCAAGAGACTG 1245
Db 1 TTCAGGCACTAAGAGGGGCTGGACCTGGCGGCGAGGAGCAAGAGACTG 50

RESULT 26

AA23425

ID AAX23425 standard; DNA; 701 BP.

XX AAX23425;

AC AAX23425;

DT 18-JUN-1999 (first entry)

XX Mouse TNRL3 DNA.

DE Mouse TNRL3 DNA.

XX Tumour necrosis factor receptor; signal transducer molecule; TNF
developmental abnormality; gestational abnormality; prostate c
APO6; APO8; APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy;
cytoplaemic domain; immunogen; antibody preparation; breast carc
apoptosis; mouse; ss.

OS Mus sp.

XX Key Location/Qualifiers

FT CDS 1..636

FT /*tag= a

FT /product= "TNRL3"

XX WO9911791-A2.

XX PD 11-MAR-1999.

XX 04-SEP-1998; 98WO-US018393.

XX 05-SEP-1997; 97US-00924634.

XX (UNIW) UNIV WASHINGTON.

XX Chaudhary PM;

XX WPI; 1999-205191/17.

XX P-PSDB; AAW93591.

XX New Tumor Necrosis Factor family receptor polypeptides and ligan
useful for diagnosis and treatment of prostate cancer and develo
or gestational abnormalities.

XX

Fig 13B; 156pp; English.

n describes isolated Tumor Necrosis Factor (TNF) family peptides: APO4, APO6, APO8 and APO9 or their active fragments. APO4 is useful for diagnosing prostate cancer by levels of APO4 in an individual. Prostate cancer can also be APO4 selective binding agents linked to a therapeutic polypeptides are also useful for identifying selective s, useful in diagnosis/treatment of disease by binding of polypeptide/active fragment which is extracellular, or the cell surface. The binding is preferably performed in lypeptides/ active fragments are also useful for screening and antagonists by binding and observing the change in APO4 active pharmacological agents useful in diagnosis or disease are also identified using APO4 polypeptides/active APO4 signal transducer molecules that specifically interact asmc domain of APO4 and detecting a change in level of APO4 method is performed in vivo or in vitro. APO polypeptides l as immunogens for preparing antibodies. APO4 is also agnosis/treatment of developmental or gestational . APO8 was transfected to human breast carcinoma cell line duced apoptosis

BP; 139 A; 210 C; 203 G; 149 T; 0 U; 0 Other;

3.4%; Score 46; DB 2; Length 701;

arity 100.0%; Pred. No. 5e-11; Indels 0; Gaps 0;

conservative 0; Mismatches 0; Indels 0; Gaps 0;

TGGGCTCTACTGCTACTGCTAGGTGCTACTTTGATGAGGG 620

TGGGCTCTACTGCTACTGCTAGGTGCTACTTTGATGAGGG 401

dard; cDNA; 1168 BP.

(first entry)

tumour necrosis factor related ligand (TRELL) gene.

necrosis factor related ligand; tnfr; treatment; cancer; sease; immune system; stimulation; suppression; on; ds.

Location/Qualifiers

2..679

/tag= a

/note= "tumour necrosis factor related ligand"

97WO-US013945.

96US-0023541P.

96US-0028515P.

97US-0040820P.

N INC.

GENEVA FACULTY MEDICINE.

e Y, Browning JL;

619/13.

524.

XX

PT Tumour necrosis factor related ligand - useful for, e.g. treating auto-immune disease and immune responses to tissue grafts.

XX Claim 2; Page 45-46; 69pp; English.

XX The sequence is that encoding mouse tumour necrosis factor related (TRELL). TRELL or active fragments can be included with a carrier pharmaceutical compositions to treat cancer, autoimmune diseases immune responses to tissue grafts, or to stimulate or suppress the system. It is useful to screen for TRELL receptors, by labelling detectable label and screening compositions for binding. Agents interfering with TRELL-receptor binding can also be screened for, then be administered, optionally with interferon- gamma, to induce death or treat, suppress or alter immune responses (especially in human adenocarcinoma cells) involving a signal pathway between the its receptor. The DNA sequence can be used in gene therapy for the related disorders in mammals (especially humans), e.g. tumours, autoimmune and inflammatory diseases or inherited genetic disorders introducing into cells, and expressing, therapeutically effective of a vector, e.g. a virus comprising a gene encoding TRELL. It may be of use in the preparation of probe probes for screening natural/synthetic DNAs for TRELL-encoding sequences and for anti-therapy

XX Sequence 1168 BP; 242 A; 360 C; 298 G; 268 T; 0 U; 0 Other;

Query Match 3.4%; Score 46; DB 2; Length 1168;

Best Local Similarity 100.0%; Pred. No. 4.9e-11;

Matches 46; Conservative 0; Mismatches 0; Indels 0; C

QY 575 GGGCTGGGCTCTACTGCTAGGTGCTACTTTGATGAGGG 620

DB 399 GGGCTGGGCTCTACTGCTAGGTGCTACTTTGATGAGGG 444

RESULT 28

ABX37032

ID ABX37032 standard; cDNA; 408 BP.

AC ABX37032;

DT 20-FEB-2003 (first entry)

DE Bovine EST associated with lactation/muscle/fat deposition #2197.

XX Bovine; ss; EST; expressed sequence tag; lactation; LMFD;

KW muscle deposition; fat deposition; genome mapping; gene identification; cattle breeding.

KW gene analysis; cattle breeding.

OS Bos Taurus.

PN US2002137139-A1.

XX 26-SEP-2002.

PF 24-SEP-2001; 2001US-00960352.

XX 12-JAN-1999; 99US-0115707P.

PR 11-JAN-2000; 2000US-00480902.

XX (BYAT/) BYATT J C.

PA (MATH/) MATHIALAGAN N.

PA (TAON/) TAO N.

PA (WARR/) WARREN W C.

XX Byatt JC, Mathialagan N, Tao N, Warren WC;

XX WPI; 2003-110599/10.

XX New nucleic acid associated with lactation, and muscle and fat deposition, useful for genome mapping, gene identification and cattle breeding, or for genetically improving cattle.

PT cattle breeding, or for genetically improving cattle.

PT cattle breeding, or for genetically improving cattle.

06:25:14 2004

us-09-245-198a-3.oligo.rng

ID NO 2197; 245pp; English.

relates to a purified nucleic acid molecule associated with muscle and fat deposition (designated LMFD), derived from the LMFD nucleic acid can specifically hybridise to a second molecule comprising any of 15112 nucleotide sequences, ABX34836-ABX49947, or complements of them. Also included are a transformed cell having a nucleic acid comprising an LMFD nucleic acid to a promoter and a 3' non-translated sequence that the cell to cause termination of transcription and addition of ribonucleotides to a 3' end of the mRNA molecule; and ing a level or pattern of a molecule in a bovine cell or using: (a) incubating a marker nucleic acid (comprising any nucleic acid sequences or its complement or fragment) with a y nucleic acid molecule obtained from the bovine cell or e hybridisation between the marker nucleic acid and the y nucleic acid permits the detection of the molecule; and (b) a level or pattern of the complementary nucleic acid, where n of the complementary nucleic acid is predictive of the term of the molecule. The LMFD nucleic acid is used for a level or pattern of a molecule in a bovine cell or tissue, for genome mapping, gene identification and analysis, cattle separation of constructs for use in cattle gene expression, or lly improving cattle. The present sequence is one of the LMFD EST (expressed sequence tag) nucleic acids. Note: The ence was not shown in the specification but was obtained in ormat from the USPTO web site: o.gov/sequence.html?DocID=20020137139

BP; 78 A; 136 C; 137 G; 57 T; 0 U; 0 Other;

2.6%; Score 36; DB 7; Length 408;

larity 100.0%; Pred. No. 2.3e-06;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GGCTGCTCGCTCGCTCGCTGGCGTGGTCAG 233

GGCTGCTCGCTCGCTCGCTGGCGTGGTCAG 152

ndard; DNA; 412 BP.

(first entry)

ated gene, SEQ ID 1352.

; gene expression profile; hepatotoxicity; liver; ng; toxicity assay; ds.

-A2.

2003WO-US003194.

2002US-00060087.

2002US-0364045P.

2002US-0364055P.

2002US-0436643P.

; LOGIC INC.

Porter M, Johnson K, Higgs B, Castle A, Elashoff M;

19530/65.

PT Predicting a toxic effect of a compound, useful in identifying t
PT markers in liver tissues or cells for drug screening and toxicit
PT comprises preparing gene expression profile of tissue or cells e
PT the compound.

XX Claim 1; SEQ ID NO 1352; 1156pp; English.

XX The present invention relates to a method for predicting a toxic
CC of a compound. The method comprises preparing a gene expression
CC of a tissue or cell sample exposed to the compound, and comparir
CC gene expression profile to a database comprising SEQ ID 1-4925;
CC differential expression of the gene indicates at least one toxic
CC The method is useful for predicting at least one toxic effect of
CC compound, predicting hepatotoxicity or the progression of a toxi
CC of a compound, identifying an agent that modulates the onset or
CC progression of a toxic response, predicting the cellular pathway
CC compound modulates in a cell, and identifying an agent that modu
CC least one activity of a protein. The method and compositions of
CC present invention using a database of genes having liver toxin-i
CC differential expression, are useful in identifying toxicity mark
CC liver tissues or cells for drug screening and toxicity assays. N
CC sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly fr
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 412 BP; 104 A; 87 C; 116 G; 105 T; 0 U; 0 Other;

Query Match 2.1%; Score 29; DB 9; Length 412;

Best Local Similarity 100.0%; Pred. No. 0.0043;

Matches 29; Conservative 0; Mismatches 0; Indels 0;

QY 1336 TATTTTATTTATTTGTGACAAAATGTT 1364

Db 67 TATTTTATTTATTTGTGACAAAATGTT 39

RESULT 30

ABK40356/c

ID ABK40356 standard; DNA; 26 BP.

XX AC ABK40356;

XX 15-JUL-2002 (first entry)

XX Reverse PCR primer for gene amplification analysis of human PRO:
XX Human; PRO; benign tumour; malignant tumour; lymphoid malignancy
XX leukaemia; neuronal disorder; stromal disorder; blastocoeleic dis
XX inflammatory disorder; immune disorder; angiogenic disorder; cyt
XX neuroprotective; PCR; primer; ss.

XX Homo sapiens.

XX WO200153486-A1.

XX 26-JUL-2001.

XX 11-FEB-2000; 2000WO-US003565.

XX 08-MAR-1999; 99WO-US005028.

XX 11-MAR-1999; 99US-0123972P.

XX 11-MAY-1999; 99US-0133459P.

XX 02-JUN-1999; 99WO-US012252.

XX 22-JUN-1999; 99US-0140650P.

XX 22-JUN-1999; 99US-0140653P.

XX 20-JUL-1999; 99US-0144758P.

XX 26-JUL-1999; 99US-0145698P.

XX 17-AUG-1999; 99US-0146222P.

XX 31-AUG-1999; 99US-0151689P.

XX 01-SEP-1999; 99WO-US020111.

XX 15-SEP-1999; 99WO-US021090.

XX 30-NOV-1999; 99WO-US028313.

99WO-US028301.
99WO-US028634.
2000WO-US000219.

TECH INC.

Goddard A, Godowski PJ, Gurney AL, Hillan KJ;
Pan J, Pitti RM, Roy MA, Smith V, Stone DM;
Wood WI;

567/26.

ucleic acids encoding PRO polypeptides, useful for treating
ignant tumors, leukemias and lymphoid malignancies,
angiogenic and immunologic disorders.

age 140; 302pp; English.

nvention relates to the isolation of novel human PRO
(AAU86128-AAU86162) and the polynucleotide sequences
. The PRO polypeptides, agonists, antagonists or anti-PRO
e useful for treating benign or malignant tumours (e.g.
, bladder, breast, etc), leukaemias and lymphoid
other disorders such as neuronal, glial, astrocytal,
glandular, macrophagal, stromal and blastocoelec disorders,
immune and angiogenic disorders. The polynucleotide
also useful in gene therapy. The present sequence
PCR primer used in the methods of the present invention

P; 7 A; 1 C; 9 G; 9 T; 0 U; 0 Other;

arity 1.9%; Score 26; DB 6; Length 26;
arity 100.0%; Pred. No. 0.12;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

TCCACATAAATACAGTATCC 1030

TCCACATAAATACAGTATCC 1

dard; DNA; 140 BP.

(first entry)

ssion-related sequence, SEQ ID 542.

irucide; apoptotic; gene therapy; tumour suppression;
ion; apoptosis; virus resistance; viral infection; tumour;
tive disease; ds.

2.

2002WO-FR000543.

2001FR-00001925.

ULAR ENGINES LAB.

Amson R, Tuijnder M, Susini L;

286/05.

cid encoding a translationally controlled tumor protein,
eating, preventing and diagnosing viral, tumor or
diseases.

XX Disclosure; Page: 45pp; French.
XX
CC The present invention relates to novel nucleic acid sequences (AE
CC ABZ79313), which are involved in the molecular pathways of tumour
CC suppression, tumour reversion, apoptosis and/or virus resistance.
CC sequences are also useful for treatment or prevention of viral, t
CC and cell degenerative diseases, and also for diagnosis and progn
CC these diseases. Note: The sequence data for this patent is not
CC represented in the printed specification but is based on sequence
CC information supplied by the European Patent Office

XX Sequence 140 BP; 43 A; 23 C; 28 G; 46 T; 0 U; 0 Other;

Query Match 1.9%; Score 26; DB 7; Length 140;
Best Local Similarity 100.0%; Pred. No. 0.11;
Matches 26; Conservative 0; Mismatches 0; Indels 0; G

Qy 1 ATGTCATTGTAGACTTTGAAATTC 26

Db 67 ATGTCATTGTAGACTTTGAAATTC 42

RESULT 32

ABZ09382/c

ID ABZ09382 standard; DNA; 140 BP.

XX AC ABZ09382;

DT 16-JAN-2003 (first entry)

XX Human oligonucleotide SEQ ID 542.

Human: tumour suppressor; virucide; cytostatic; nootropic;
neuroprotective; neuroleptic; gene therapy; tumour suppression;
tumour reversion; apoptosis; viral resistance; viral infection;
cell degeneration; Alzheimer's disease; schizophrenia; cancer; ds

XX Homo sapiens.

XX FR2822475-A1.

XX 27-SEP-2002.

XX 20-MAR-2002; 2002FR-00003459.

XX 13-FEB-2001; 2001FR-00001925.

XX (MOLE-) MOLECULAR ENGINES LAB SA.

XX Telerman A, Amson R, Tuijnder M, Susini L;

XX WPI; 2003-032204/03.

XX New human nucleic acid, useful for diagnosis, prognosis and treat
e.g. of tumors, also related vectors, transformed cell, polypepti
antibodies.

XX Disclosure; Page 120; 189pp; French.

XX The present invention relates to human oligonucleotides (ABZ08841
CC ABZ09860). The expression of the oligonucleotides is implicated i
CC suppression or reversion, apoptosis and/or viral resistance. The
CC oligonucleotides are useful for preventing and/or treating viral
CC infection, tumour development and cell degeneration (e.g. Alzheim
CC disease and schizophrenia), especially cancer

XX Sequence 140 BP; 43 A; 23 C; 28 G; 46 T; 0 U; 0 Other;

Query Match 1.9%; Score 26; DB 7; Length 140;
Best Local Similarity 100.0%; Pred. No. 0.11;
Matches 26; Conservative 0; Mismatches 0; Indels 0; G

06:25:14 2004

us-09-245-198a-3.oligo.rng

TCATTGTTAGACTTTGAAATTC 26
|||||
TCATTGTTAGACTTTGAAATTC 42

ndard; DNA; 145 BP.

(first entry)

ession-related sequence, SEQ ID 15.

virucide; apoptotic; gene therapy; tumour suppression;
sion; apoptosis; virus resistance; viral infection; tumour;
ative disease; ds.

A2.

2002WO-FR000543.

2001FR-00001925.

CULAR ENGINES LAB.

Amson R, Tuijnder M, Susini L;

9286/05.

acid encoding a translationally controlled tumor protein,
reating, preventing and diagnosing viral, tumor or
diseases.

Page; 45pp; French.

invention relates to novel nucleic acid sequences (ABZ78294-
hich are involved in the molecular pathways of tumour
tumour reversion, apoptosis and/or virus resistance. The
e also useful for treatment or prevention of viral, tumour
enerative diseases, and also for diagnosis and prognosis of
es. Note: The sequence data for this patent is not
in the printed specification but is based on sequence
supplied by the European Patent Office

BP; 51 A; 28 C; 23 G; 43 T; 0 U; 0 Other;

1.9%; Score 26; DB 7; Length 145;
larity 100.0%; Pred. No. 0.11;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

TCATTGTTAGACTTTGAAATTC 26
|||||
TCATTGTTAGACTTTGAAATTC 99

ndard; DNA; 145 BP.

(first entry)

ession-related sequence, SEQ ID 135.

virucide; apoptotic; gene therapy; tumour suppression;
sion; apoptosis; virus resistance; viral infection; tumour;

KW cell degenerative disease; ds.

XX Unidentified.

OS WO200264731-A2.

PN 22-AUG-2002.

XX 13-FEB-2002; 2002WO-FR000543.

XX 13-FEB-2001; 2001FR-00001925.

PR (MOLE-) MOLECULAR ENGINES LAB.

PA Telerman A, Amson R, Tuijnder M, Susini L;

XX WPI; 2003-058286/05.

DR New nucleic acid encoding a translationally controlled tumor pro
PT useful for treating, preventing and diagnosing viral, tumor or
PT degenerative diseases.

XX Disclosure; Page; 45pp; French.

XX The present invention relates to novel nucleic acid sequences (A
CC ABZ79313), which are involved in the molecular pathways of tumou
CC suppression, tumour reversion, apoptosis and/or virus resistance
CC sequences are also useful for treatment or prevention of viral,
CC and cell degenerative diseases, and also for diagnosis and progn
CC these diseases. Note: The sequence data for this patent is not
CC represented in the printed specification but is based on sequenc
CC information supplied by the European Patent Office

XX Sequence 145 BP; 51 A; 28 C; 23 G; 43 T; 0 U; 0 Other;

Query Match 1.9%; Score 26; DB 7; Length 145;
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Db 74 ATGTCATTGTTAGACTTTGAAATTC 99

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XX AC ABZ08955;

XX 16-JAN-2003 (first entry)

XX Human oligonucleotide SEQ ID 15.

XX Human; tumour suppressor; virucide; cytostatic; nootropic;
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KW tumour reversion; apoptosis; viral resistance; viral infection;
KW cell degeneration; Alzheimer's disease; schizophrenia; cancer; d
XX Homo sapiens.

XX FR2822475-A1.

XX 27-SEP-2002.

XX 20-MAR-2002; 2002FR-00003459.

XX 13-FEB-2001; 2001FR-00001925.

XX (MOLE-) MOLECULAR ENGINES LAB SA.

XX Telerman A, Amson R, Tuijnder M, Susini L;

XX

204/03.
leic acid, useful for diagnosis, prognosis and treatment.
3, also related vectors, transformed cell, polypeptides and
age 40; 189pp; French.
vention relates to human oligonucleotides (ABZ08841-
e expression of the oligonucleotides is implicated in tumour
r reversion, apoptosis and/or viral resistance. The
des are useful for preventing and/or treating viral
mour development and cell degeneration (e.g. Alzheimer's
chizophrenia), especially cancer
BP; 51 A; 28 C; 23 G; 43 T; 0 U; 0 Other;
arity 1.9%; Score 26; DB 7; Length 145;
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CATTGTTAGACTTTGAAATTC 99
dard; DNA; 145 BP.
(first entry)
cleotide SEQ ID 135.
suppressor; virucide; cytostatic; nootropic;
ve; neuroleptic; gene therapy; tumour suppression;
ion; apoptosis; viral resistance; viral infection;
tion; Alzheimer's disease; schizophrenia; cancer; ds.
2002FR-00003459.
2001FR-00001925.
JLAR ENGINES LAB SA.
Amson R, Tuijnder M, Susini L;
204/03.
leic acid, useful for diagnosis, prognosis and treatment.
3, also related vectors, transformed cell, polypeptides and
age 58; 189pp; French.
vention relates to human oligonucleotides (ABZ08841-
e expression of the oligonucleotides is implicated in tumour
r reversion, apoptosis and/or viral resistance. The
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chizophrenia), especially cancer
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XX OS Homo sapiens.
XX PN WO9514772-A1.
XX PD 01-JUN-1995.
XX PF 11-NOV-1994; 94WO-JP001916.
XX PR 12-NOV-1993; 93JP-00355504.
XX PA (MATS/) MATSUBARA K.
XX PA (OKUB/) OKUBO K.
XX PI Matsubara K, Okubo K;
XX WI; 1995-206931/27.
Single-stranded DNA for identifying gene signatures - isolated fr
directed human cDNA library that reflects relative abundance of c
mRNA in specific human tissues.
Claim 1; Page 456; 2245pp; Japanese.
A single-stranded DNA (or its complementary strand or the corresp
-stranded DNA) which comprises one of the 7837 "GS" sequences giv
AAT19001-R26837 and which is able to hybridise to part of human g
DNA, cDNA or mRNA is claimed. The GS (Gene Signature) sequences w
obtained from 3'-directed cDNA libraries prepared from various hu
tissues; synthesis of cDNA was initiated from the 3'-end of mRNA
poly(T) as the sole primer. Since the 3'- untranslated sequence i
to a particular mRNA species, almost all the 3'-oriented cDNAs hy
with specific mRNAs. Each library is constructed so as to reflect
accurately the relative abundance of different mRNAs in the parti
tissue from which it was derived. The appearance frequency of a g
in a cDNA library can be determined (esp. using primers and probe
CC derived from the GS sequences) as a means of diagnosing abnormal
CC function or for recognising different cell types
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Vogt, J.L., Wetherby, K.D., Wiggins, L., Young, A. and Green,
NISC Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 218485)
Green, E.D.
Direct Submission
Submitted (17-JUN-2002) NIH Intramural Sequencing Center,
Grovmont Circle, Gaithersburg, MD 20877, USA
3 (bases 1 to 218485)
Green, E.D.
Direct Submission
Submitted (05-JUN-2003) NIH Intramural Sequencing Center,
Grovmont Circle, Gaithersburg, MD 20877, USA
On Jun 5, 2003 this sequence version replaced gi:26449071
----- Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: <http://www.nisc.nih.gov>
Contact: nisc_zoo@nih.gov
----- Project Information
Center project name: cms
Center clone name: 145D13

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicated order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones, alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.90319
Consensus quality: 214085 bases at least Q40
Consensus quality: 215055 bases at least Q30
Consensus quality: 216264 bases at least Q20
Insert size: 190000; agarose-ftp
Insert size: 216885; sum-of-contigs
Quality coverage: 12.65x in Q20 bases; agarose-ftp
Quality coverage: 11.08x in Q20 bases; sum-of-contigs

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* NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that ha
* ve provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
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* 7549 7648: gap of unknown length
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* 31918 32017: gap of unknown length
* 32018 50433: contig of 18416 bp in length
* 50434 50533: gap of unknown length
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* 128726 143521: contig of 14796 bp in length

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1998

Submission
 :ed (30-JUL-2003) Cell Biology, Biogen, 12 Cambridge Center,
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 ice update by submitter
 - 30, 2003 this sequence version replaced gi:2707220.
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 Wiley, S.R.
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 JOURNAL
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 IMMUNEX CORPORATION (US)
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1.4	493	8	ACH34013	Ach34013 Human end
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1.9	282	2	AAT22190	Aat22190 Human gen
1.1	195	6	ABK29540	Abk29540 Colon ade
1.1	412	9	ADB56326	Adb56326 Toxicity
1.7	264	7	ABX52254	Abx52254 Bovine ES
1.1	114955	2	AAV53491	Aax53491 Human ade
1.9	3163	9	ADC87060	Adc87060 Human GPC

24	65.4	4.8	1064	6	ABT09678	Hu
C 25	63.8	4.6	5452	9	ADC86736	Hu
C 26	62.2	4.5	12733	6	ABK98631	Ve
C 27	62.2	4.5	12733	8	ACD13882	L.
C 28	62.2	4.5	12739	6	ABK98592	Ve
C 29	62.2	4.5	12739	8	ACD13843	Fl
C 30	61.4	4.5	1117	9	ADC86688	Hu
C 31	61	4.4	1337	2	AAZ17263	Hu
C 32	60.6	4.4	1000	3	AAA02484	Hu
C 33	60	4.4	60	6	ABN58848	Hu
C 34	60	4.4	60	6	ABN41049	Hu
C 35	60	4.4	60	6	ABN58593	Hu
C 36	60	4.4	60	6	ABN58591	Hu
C 37	60	4.4	60	6	ABN58849	Hu
C 38	59.6	4.3	3133	9	ADC86738	Hu
C 39	59.2	4.3	10732	3	AAA10594	Ge
C 40	58	4.2	1218	3	AAA02488	Hu
C 41	56	4.1	1065	6	ABT09682	Hu
C 42	54.8	4.0	1017	7	ADJ36876	S.
C 43	54.8	4.0	2000	7	ADA71938	Ri
C 44	54.8	4.0	29870	7	ADJ36874	St
C 45	54.4	4.0	600	6	ABQ52497	Ol

ALIGNMENTS

RESULT 1
AAV18600
ID AAV18600 standard; cDNA; 1373 BP.
XX AC AAV18600;
XX AC AAV18600;
DT 21-JUL-1998 (first entry)
XX Homo sapiens tumour necrosis factor related ligand (TRELL) gene.
DE TRELL; tumour necrosis factor related ligand; tnf; treatment; can
KW autoimmune disease; immune system; stimulation; suppression;
KW graft rejection; ds.
XX Homo sapiens.
XX Key Location/Qualifiers
FH 1. 852
FT CDS /tag= a
FT /note= "tumour necrosis factor related ligand"
XX W09805783-A1.
XX 12-FEB-1998.
XX 07-AUG-1997; 97WO-US013945.
XX 07-AUG-1996; 96US-0023541P.
XX 18-OCT-1996; 96US-0028515P.
XX 18-MAR-1997; 97US-0040820P.
XX (BIOJ) BIOGEN INC.
XX (UYGE-) UNIV GENEVA FACULTY MEDICINE.
XX Chicheportiche Y, Browning JL;
XX WPI: 1998-145619/13.
XX P-PSDB; AAW47525.
XX Tumour necrosis factor related ligand - useful for, e.g. treating
XX auto-immune disease and immune responses to tissue grafts.
XX Claim 2; Page 48-50; 69pp; English.
XX The sequence is that encoding human tumour necrosis factor relate
XX (TRELL). TRELL or active fragments can be included with a carrier
CC

XX
PA (GEMY) GENETICS INST INC.

92...841
/*tag= a
/product= "Apo-3 ligand"

98WO-US021407.

97US-0062037P.

97US-0069862P.

VTech INC.

, Marsters SA, Pitti R;

7982/24.

9369.

3- ligand (a tumor necrosis factor) homologue.

31: 74pp; English.

sequence encodes a human tumour necrosis factor (TNF) and homologue designated Apo-3 ligand. Apo-3 ligand has activity. Apo-3 ligand can be used to induce apoptosis in cancer cells, to induce NF-kappaB-dependent transcription and /SAPK-dependent responses in mammalian cells

1 BP; 281 A; 464 C; 404 G; 272 T; 0 U; 0 Other;

96.5%; Score 1325; DB 2; Length 1421;

arity 98.5%; Pred.No. 3.1e-272;

conservative 0; Mismatches 20; Indels 0; Gaps 0;

AAATTTCCGGCCCGCGCTCCCTCCCGGATCCCTCGGTCCTCGGATCGGGGG 76

AGATCCCTCGACCTCGACCCACCGCTCCCGATCCCTCGGTCCTCGGATCGGGGG 62

TGAGGAGGACAGAGCCCGCCCATGGCCCGCCCGTCCGAGCCAGAGCGGAGGG 136

TGAGGAGGACAGAGCCCGCCCATGGCCCGCCCGTCCGAGCCAGAGCGGAGGG 122

TCCGGGGAGCGGGACCGCCCTGCTGCTGCCCTCGCTCGGTCCTCGGTCCTGG 196

TCCGGGGAGCGGGACCGCCCTGCTGCTGCCCTCGCTCGGTCCTCGGTCCTGG 182

TGGCTCGCTCGGTCCTCGCTGCTGCCCTCGCTGCTGCCCTCGCTGCTGCCCT 256

TGGCTCGCTCGGTCCTCGCTGCTGCCCTCGCTGCTGCCCTCGCTGCTGCCCT 242

TCCAGGAGCTCGCCAGGAGGAGCTGGTGGCAGAGGAGGACAGAGCCCGTCCGAAC 316

TCCAGGAGCTCGCCAGGAGGAGCTGGTGGCAGAGGAGGACAGAGCCCGTCCGAAC 302

TCCCCAGACAGAGAGAGCCAGAGTCCCTGGCTTCCCTGAAACCGACTAGTTCCGC 376

TCCCCAGACAGAGAGAGCCAGAGTCCCTGGCTTCCCTGAAACCGACTAGTTCCGC 362

TCCAGAGTCCCTGAAAGCCGGAACACCGGCTCGAGAGCGGATCGAGCCCAT 436

TCCAGAGTCCCTGAAAGCCGGAACACCGGCTCGAGAGCGGATCGAGCCCAT 422

TAGTTTCATCCAGCTGACAGAGAGGAGCGGATGTTGGAGCGGACAGTGA 496

TAGTTTCATCCAGCTGACAGAGAGGAGCGGATGTTGGAGCGGACAGTGA 482

TCTGGAGGAGCCAGAGTCAACAGCTCCAGCCCTCTGCGCTACAAACCGCCAGATCG 556

TCTGGAGGAGCCAGAGTCAACAGCTCCAGCCCTCTGCGCTACAAACCGCCAGATCG 542

TGTTTATAGTCACCCGGGCTGGGCTCTACTACTGTCTAGGTGCATTTGATG 616

Db 543 GGGAGTTTATAGTCACCCGGGCTGGGCTCTACTACTGTCTACTGTCTAGGTGCATTT
QY 617 AGGGGAAGGCTGTCTACTTGAAGCTGGAGCTTCTGCTGGTGGATGGTGGCCCTT
Db 603 AGGGGAAGGCTGTCTACTTGAAGCTGGAGCTTCTGCTGGTGGATGGTGGCCCTT
QY 677 GCCTGGAGGAATTTCTCAGCACTGGGGCCAGTTCCCTCGGGCCCGCAGCTCCGCTT
Db 663 GCCTGGAGGAATTTCTCAGCACTGGGGCCAGTTCCCTCGGGCCCGCAGCTCCGCTT
QY 737 AGGTGTCTGGGCTGTGGCCCTGGGGCCAGGGTCTCTCTCTGGGATCCGACCCCTT
Db 723 AGGTGTCTGGGCTGTGGCCCTGGGGCCAGGGTCTCTCTCTGGGATCCGACCCCTT
QY 797 GGGCCCATCTCAAGGCTGCCCTTCTCACTCACTCTCGGACTCTTCAGGTTTCA
Db 783 GGGCCCATCTCAAGGCTGCCCTTCTCACTCACTCTCGGACTCTTCAGGTTTCA
QY 857 GGGCCCTGTCTCTCCCACTGGTCTCCAGGCTGCCGGTCTCCCTCGACAGCTCTCT
Db 843 GGGCCCTGTCTCTCCCACTGGTCTCCAGGCTGCCGGTCTCCCTCGACAGCTCTCT
QY 917 ACCCGTCTCTCTCCCACTGGTCTCCAGGCTGCCGGTCTCTCTCTCGACAGCTCTCT
Db 903 ACCCGTCTCTCTCCCACTGGTCTCCAGGCTGCCGGTCTCTCTCTCGACAGCTCTCT
QY 977 GAGGCTGTCTGGGCTGTCTCAGCTGTCTTCATCCACATAAATACAGTATTTCCC
Db 963 GAGGCTGTCTGGGCTGTCTCAGCTGTCTTCATCCACATAAATACAGTATTTCCC
QY 1037 TATCTTCAAACTCCCACTGGGCTCTCCAGCTCTCCAGCTCTCTCCCAATCCCTG
Db 1023 TATCTTCAAACTCCCACTGGGCTCTCCAGCTCTCCAGCTCTCTCCCAATCCCTG
QY 1097 TTGAGGCTCTGGGCTGTCTCAGCTCTCCCTGGGCTCCAGCTCTCCAGCTCTGT
Db 1083 TTGAGGCTCTGGGCTGTCTCAGCTCTCCCTGGGCTCCAGCTCTCCAGCTCTGT
QY 1157 CTGTACTCTGTGGGCAAGATGGGTCCAGAGACCCCTTCAGGCTCTCCAGCTCTGT
Db 1143 CTGTACTCTGTGGGCAAGATGGGTCCAGAGACCCCTTCAGGCTCTCCAGCTCTGT
QY 1217 GACCTGGGCGGAGGAGCCAAAGAGCTGGGCTTAGGCGGAGGTTCCCAATGT
Db 1203 GACCTGGGCGGAGGAGCCAAAGAGCTGGGCTTAGGCGGAGGTTCCCAATGT
QY 1277 GCGAGAAACAGACAGCTCTCTCTTCCAGATTCCTGTGGATTTTAAACAG
Db 1263 GCGAGAAACAGACAGCTCTCTCTTCCAGATTCCTGTGGATTTTAAACAG
QY 1337 ATTTTATTTATTTATTTGTGACAAATGTTGATAAATGG 1373
Db 1323 ATTTTATTTATTTATTTGTGACAAATGTTGATAAATGG 1359

RESULT 4

AAA49717

ID AAA49717 standard; cDNA; 1353 BP.

XX AC AAA49717;

XX XX 25-SEP-2000 (first entry)

XX DE Human PRO207 cDNA clone DNA30879-1152.

XX DE PRO207; human; antitumor; tumour; therapy; cytostatic; breast ca;

XX KW ovarian cancer; renal cancer; colorectal cancer; uterine cancer;

XX KW prostate cancer; lung cancer; bladder cancer;

XX KW central nervous system cancer; melanoma; leukaemia; neoplasm; ss.

XX OS Homo sapiens.

XX XX

Location/Qualifiers

58..807
/*tag= a
58..177
/*tag= b
178..804
/*tag= c

2.

99WO-US028565.

98US-0113296P.
99WO-US005028.
99US-0130232P.
99US-0131445P.
99US-0134287P.
99US-0144758P.
99US-0145698P.
99WO-US021090.
99WO-US021547.

TECH INC.

Goddard A, Godowski PJ, Gurney AL, Marsters SA;
tbi RM, Wood WI;

588/38.
138.

ion to inhibit neoplastic cell growth or for treating tumor
risers polypeptides PRO179, PRO207, PRO320, PRO219, PRO221,
1, PRO301, PRO526, PRO362, PRO356, PRO509 or PRO866.

3; 172pp; English.

quence is that of cDNA clone DNA30879-1152 (ATCC 209358)
PRO207 (see AAY95338), which shows homology to several
tumour necrosis factor family, especially human
3.4%). The cDNA was identified in a foetal kidney cDNA
ing identification of an expressed sequence tag with
man Apo-2 ligand. A claimed method for inhibiting the
mour cell comprises exposing the tumor cell to PRO179,
PRO219, PRO221, PRO224, PRO328, PRO301, PRO526, PRO362,
or PRO866 (see AAY95337-49), their agonists or chimeric
incorporating them. The tumour is especially a cancer
breast, ovarian, renal, colorectal, uterine, prostate,
and central nervous system cancer, melanoma and leukaemia.
encoding PRO179 etc. are used in the recombinant production
our polypeptides

BP; 257 A; 443 C; 389 G; 264 T; 0 U; 0 Other;

96.2%; Score 1320.2; DB 3; Length 1353;
arity 99.8%; Pred. No. 3.3e-271;
nservative 0; Mismatches 3; Indels 0; Gaps 0;

CTCTGGTCTCCGGGATGGGGGCGGTAGGAGCAGACAGCCCCCGCCCCCATG 108
CTCTGGGTCCTGGGATGGGGGCGGTAGGAGCAGACAGCCCCCGCCCCCATG 60

CCGTGGAGCCAGAGCGGAGGGGGCGCGGGGGGAGCGGGCACCGCCCTGCTG 168
CCGTGGAGCCAGAGCGGAGGGGGCGCGGGGGGAGCGGGCACCGCCCTGCTG 120

CTCTGGTCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGG 228
CTCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGGGCTGG 180

TTTGGGAGCCGGGATCGCTCTGGCCCGAGGAGCTGCCAGGAGGAGCTGGTG 288
TTTGGGAGCCGGGATCGCTCTGGCCCGAGGAGCTGCCAGGAGGAGCTGGTG

181 GTCAGTTTGGGGAGCCGGGCGATCGCTGTCCGCGCCAGGAGCCTGCCAGGAGGAGCT
289 GCAGAGGAGGACCCAGGACCGTCGGAACCTGAATCCCCAGACAGAAAGAACCCAGGA
241 GCAGAGGAGGACCCAGGACCGTCGGAACCTGAATCCCCAGACAGAAAGAACCCAGGA
349 GCGCCTTTCTGTAAACCGACTAGTTTCGGCCTCGCAGAGTGCACCTAAAGGCCGGA
301 GCGCCTTTCTGTAAACCGACTAGTTTCGGCCTCGCAGAGTGCACCTAAAGGCCGGA
409 CGGGCTCGAAGAGCGATCGAGCCCAATTATGAAGTTTATCCACGACCTGACAGGA
361 CGGGCTCGAAGAGCGATCGAGCCCAATTATGAAGTTTATCCACGACCTGACAGGA
469 GCGCAGCAGGTGTGACAGCGGACAGTGAGTGGCTGGGAGGAGCCAGAAATCAACAG
421 GCGCAGCAGGTGTGACAGCGGACAGTGAGTGGCTGGGAGGAGCCAGAAATCAACAG
529 AGCCCTCTGCGCTACAAACCGCCAGATCGGGGAGTTTTATAGTCAACCGGGCTGGGCT
481 AGCCCTCTGCGCTACAAACCGCCAGATCGGGGAGTTTTATAGTCAACCGGGCTGGGCT
589 TACCTGTACTGTGAGTGCACCTTTGATGAGGAGGAGGCTGTCTACCTGAAGCTGGA
541 TACCTGTACTGTGAGTGCACCTTTGATGAGGAGGAGGCTGTCTACCTGAAGCTGGA
649 CTGGTGGATGGTGTGCTGGCCCTGCGCTGCTGAGGAGAAATTCACGACCTGCGGC
601 CTGGTGGATGGTGTGCTGGCCCTGCGCTGCTGAGGAGAAATTCACGACCTGCGGC
709 TCCTTCGGGCCCCAGCTCCGCTCTCCAGGTTCTGGGCTGTGGGCTGTGGCCCTGCGGC
661 TCCTTCGGGCCCCAGCTCCGCTCTCCAGGTTCTGGGCTGTGGGCTGTGGCCCTGCGGC
769 TCCTTCGGGCCCCAGCTCCGCTCTCCAGGTTCTGGGCTGTGGGCTGTGGCCCTGCGGC
721 TCCTTCGGGCCCCAGCTCCGCTCTCCAGGTTCTGGGCTGTGGGCTGTGGCCCTGCGGC
829 TACTTCGGACTCTTCCAGGTTCTACTGAGGGGCGCTGTCTCCCCACAGTGGTCCCA
781 TACTTCGGACTCTTCCAGGTTCTACTGAGGGGCGCTGTCTCCCCAGTGGTCCCA
889 GCGGCTCCCTCGACAGCTCTCTGGGACCCCGGTCCCTCTGCCCCACCCCTCAGC
841 GCGGCTCCCTCGACAGCTCTCTGGGACCCCGGTCCCTCTGCCCCACCCCTCAGC
949 CTTTGTCTCAGACCTCGCCCTCCCTCTAGAGGCTGCTGGGCTGTTCAGTGTCTT
901 CTTTGTCTCAGACCTCGCCCTCCCTCTAGAGGCTGCTGGGCTGTTCAGTGTCTT
1009 TCCACATAAATACAGTATTCCACCTCTTATCTTACAACTCCCCACCGCCACTC
961 TCCACATAAATACAGTATTCCACCTCTTATCTTACAACTCCCCACCGCCACTC
1069 CTTCACTAGTCCCAATCCCTGACCCCTTTGAGGCCCCCAGTGATCTGACTCCCCC
1021 CTTCACTAGTCCCAATCCCTGACCCCTTTGAGGCCCCCAGTGATCTGACTCCCCC
1129 GCCACAGCCCCCAGGCAATGTGTTCATCTACTGTGGGCAAGATGGGTCCA
1081 GCCACAGCCCCCAGGCAATGTGTTCATCTACTGTGGGCAAGATGGGTCCA
1189 ACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGGGCGAGAGCAAGAGAGACTT
1141 ACCCCACTTCAGGCACTAAGAGGGGCTGACCTGGGGCGAGAGCAAGAGAGACTT
1249 CTAGGCCAGAGTTCCTCAATGTGAGGGGCGAGAAACAAGAAAGCTCTCCCTTG
1201 CTAGGCCAGAGTTCCTCAATGTGAGGGGCGAGAAACAAGAAAGCTCTCCCTTG
1309 TTCCCTGTGATTTTAAACAGATATTTTATTTATTTATTTATTTATTTATTTATTT
1261 TTCCCTGTGATTTTAAACAGATATTTATTTATTTATTTATTTATTTATTTATTTATTT

ATCTGACTCCCCCTGGCCACAGACCCCGAGGCGATTGTTCACTGTTACTCTGTG 1080

AAGGATGGGTCCAGAGAACCCTTCCAGGCACCTTAAGAGGGGCTGGACCTGGCGGCA 1228

AAGGATGGGTCCAGAGAACCCTTCCAGGCACCTTAAGAGGGGCTGGACCTGGCGGCA 1140

AGCCAAAGAGACTGGGCGCTAGGCCAGAGATTCCCAATGTGAGGGGCGAGAAACAAG 1288

AGCCAAAGAGACTGGGCGCTAGGCCAGAGATTCCCAATGTGAGGGGCGAGAAACAAG 1200

AGCTCTCCCTCTGAGAAATCCCTGTGATTTTAAAAACAGATATTATTTTATTATT 1348

AGCTCTCCCTCTGAGAAATCCCTGTGATTTTAAAAACAGATATTATTTTATTATT 1260

GTGACAAAATGTTGATAAAATGG 1373

GTGACAAAATGTTGATAAAATGG 1285

Standard; cDNA: 1306 BP.

(first entry)

coding sequence.

; tumour necrosis factor; ligand; cytostatic;
tor; osteopathic; gene; ss.

Location/Qualifiers

18. 767

$$/\star \text{taq} = \bar{\alpha}$$

```

/ cag= a
/product= "Human TWEAK"

```

-A2.

2002WO-US023782.

2001US-0307838P.

J GENOME SCI INC.

Rosen CA:

1659/40.

1659/40
2315.

oligomeric complex having a first polypeptide member of the TNF family, a second polypeptide member of the TNF family, and a third polypeptide member of the TNF family, useful for treating cancer, osteoporosis or an autoimmune disease.

page 367-368: 388pp: English.

sequence is that of a polynucleotide encoding human TNFAIP3. This relates to compositions comprising heterotrimeric complexes of TNF α and TNF β and TNF γ ligand family members, and their use in the prevention and treatment of disease. In one embodiment, the heterotrimeric complex comprises full-length or extracellular portions of TNF α , TNF β , and TNF γ members, preferably VEGI or VEGI-SV. The heterotrimeric complex and the invention are useful for treating an autoimmune disease, such as rheumatoid arthritis, psoriasis, and particularly for inhibiting cancer cell proliferation, increasing B cell proliferation, or inducing apoptosis of cancer cells.

CACGGCCCACTCTCCACTCACTAGCTCCCAATCCCTTGACCCCTTTGAGGCCCCCA	1020
TCTTCGACTCCCCCTCTGGCCACAGACCCCCAGGCAATTGTGTCACTACTCTGTG	1168
TCTTCGACTCCCCCTCTGGCCACAGACCCCCAGGCAATTGTGTCACTACTCTGTG	1080
AGGATGGGTTCCAGAAAGACCCCACTTCAGGCACTAAGAGGGGCTGCACCTGGCGGCA	1228
AGATGGGTTCCAGAAAGACCCCACTTCAGGCACTAAGAGGGGCTGCACCTGGCGGCA	1140
GCACAAAGAGACTGGCGCTAGCGCCAGGAGTTCCTCAAAATGTGAGGGCGAGAAACAAG	1288
GCACAAAGAGACTGGCGCTAGCGCCAGGAGTTCCTCAAAATGTGAGGGCGAGAAACAAG	1200
GCTCTCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATTATTTTATTATT	1348
GCTCTCTCCCTTGAGAAATTCCTGTGGATTTTAAACAGATATTATTTTATTATT	1260
TGACAAAAATGTTGATAAATGG	1373
TGACAAAAATGTTGATAAATGG	1285

lard; cDNA; 1306 BP.

(first entry)

coding TNF ligand family member #12.

an; tumour necrosis factor; TNF ligand; endokine alpha;
a resorption disorder; osteoporosis; Paget's disease;
ification.

41.

2002US-00218547.

2001US-0312542P.

2001US-0330761P.

C A.

LI B.

Rosen CA, Nardelli B;

072/66.

906

alpha gene useful for preparing a composition for treating a patient with excessive or insufficient bone resorption e.g., Paget's disease or arterial calcification.

3Q ID NO 23; 145pp; English.

relates to an isolated nucleic acid molecule encoding a factor family ligand. A composition comprising the body or its fragment is used for treating an individual in need level of endokine alpha activity. The endokine alpha present in a heterotrimeric complex is used for treating an disorder associated with excessive bone resorption, osteoporosis, Paget's disease or arterial calcification. Treating an disorder associated with insufficient bone resorption using an endokine alpha antagonist, which is the

CCAGCGCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCTTTGAGGCCCCCA	1108
CCAGCGCCACTCTCCACCTCACTAGCTCCCAATCCCTGACCTTTGAGGCCCCCA	1020
ATCTCGACTCCCCCTGGCCACAGACCCCCAGGGCATTGTCTTCACCTGTACTCTGTG	1168
ATCTCGACTCCCCCTGGCCACAGACCCCCAGGGCATTGTCTTCACCTGTACTCTGTG	1080
AAGATGGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGCTGGACCTGGCGGCCA	1228
AAGATGGGTCCAGAGACCCCACTTCAGGCACCTAAGAGGGCTGGACCTGGCGGCCA	1140
AGCCAAAGAGACTGGGCTAGGCCAGAGATTCCCAAAATGTGAGGGCGAGAAACAAG	1288
AGCCAAAGAGACTGGGCTAGGCCAGAGATTCCCAAAATGTGAGGGCGAGAAACAAG	1200
AGCTTCCTCCCTTGAGAAATCCCTGTGGATTTTAAAAACAGATATATTTTATTATT	1348
AGCTTCCTCCCTTGAGAAATCCCTGTGGATTTTAAAAACAGATATATTTTATTATT	1260
GTGACAAAATGTTGATAAATGG	1373
GTGACAAAATGTTGATAAATGG	1285

standard; cDNA; 1236 BP.

(first entry)

endothelium proliferative agent gene.

othelium proliferative agent; TREPA; wound healing; cancer;
ing; vascularisation; apoptosis; autoimmune; birth control.

```
Location/Qualifiers
1. .750
/*tag= a
/product= "TREPA"
```

98WO-US002859.

97US-00798692.
98US-00021706.

IT LAB.

255/38.
1745.

oleic acid encoding TREPA - useful for diagnosis and autoimmune disease, tumours and inflammation.

je 123-4; 142pp; English.

ed endothelium proliferative agent (TREPA), or its agonists, are used to treat a deficit of TREPA, e.g. to healing or tissue grafting, by promoting vascularization, be apoptosis for treating cancer and eliminating autoreactive, an adjunct to cancer chemotherapy or antiviral treatment. S can also be used to target cytotoxic agents or for

100CACATAAATACAGTATTTCCACCTTTATCTTATCAACTCCCCACCGCCACTCT 1065
100CACATAAATACAGTATTTCCACCTTTATCTTATCAACTCCCCACCGCCACTCT 960
CTCAGTCTCCCAATCCCTGACCTTTGAGGCCCCCAGTGTCTGACTCCCCC 1125
CTCAGTCTCCCAATCCCTGACCTTTGAGGCCCCCAGTGTCTGACTCCCCC 1020
CCACAGCCCCCAGGCAATTTGTTCACTGTACTCTGTGGCAAGGATGGGTCCAG 1185
CCACAGCCCCCAGGCAATTTGTTCACTGTACTCTGTGGCAAGGATGGGTCCAG 1080
ACCCACTTCAGGCACTAAGAGGGGTGGACCTTGGGGGAGGAGCAAGAGACTG 1245
ACCCACTTCAGGCACTAAGAGGGGTGGACCTTGGGGGAGGAGCAAGAGACTG 1140
TAGGCGAGAGTTCCCAATGTAGGGGGGAGGAGCAAGAGCTCTCCCTTGA 1305
TAGGCGAGAGTTCCCAATGTAGGGGGGAGGAGCAAGAGCTCTCCCTTGA 1200
TCCCTGTGGATTTTAAACAGATATTTT 1341
TCCCTGTGGATTTTAAACAGATATTTT 1236

idard; DNA; 1030 BP.

(first entry)

JNA.

sis factor receptor; signal transducer molecule; TNF; APO4;
abnormality; gestational abnormality; prostate cancer;
APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy; disease;
lomain; immunogen; antibody preparation; breast carcinoma;
lman; ss.

Location/Qualifiers
1. .627
/*tag= a
/product= "TNRL3"

98WO-US018393.

97US-00924634.

WASHINGTON.

191/17.
.590.

rosis Factor family receptor polypeptides and ligands -
agnosis and treatment of prostate cancer and developmental
l abnormalities.

Fig 13A; 156pp; English.

n describes isolated Tumor Necrosis Factor (TNF) family
Peptides: APO4, APO6, APO8 and APO9 or their active
id isolated TNF related ligands 1 and 3 (TNRL1 and TNRL3) or

CC their active fragments. APO4 is useful for diagnosing prostate c
CC determining levels of APO4 in an individual. Prostate cancer can
CC treated using APO4 selective binding agents linked to a therapeu
CC moiety. APO4 polypeptides are also useful for identifying select
CC binding agents, useful in diagnosis/treatment of disease by bind
CC agents to the polypeptide/active fragment which is extracellular
CC expressed on the cell surface. The binding is preferably perform
CC vivo. APO4 polypeptides/ active fragments are also useful for sc
CC for agonists and antagonists by binding and observing the change
CC activity. Effective pharmacological agents useful in diagnosis o
CC treatment of disease are also identified using APO4 polypeptides
CC fragments and APO4 signal transducer molecules that specifically
CC with a cytoplasmic domain of APO4 and detecting a change in leve
CC activity. The method is performed in vivo or in vitro. APO polyp
CC are all useful as immunogens for preparing antibodies. APO4 is a
CC useful for diagnosis/treatment of developmental or gestational
CC abnormalities. APO8 was transfected to human breast carcinoma ce
CC MCF-7, and induced apoptosis
XX

SQ Sequence 1030 BP; 223 A; 317 C; 279 G; 211 T; 0 U; 0 Other;

Query Match 60.7%; Score 833.4; DB 2; Length 1030;
Best Local Similarity 99.9%; Pred. No. 1.1e-167;
Matches 834; Conservative 0; Mismatches 1; Indels 0;

QY 229 GTCACTTTGGGAGCCGGGCATCGCTGTCGCGCCAGGAGCCTGCCAGGAGGC
DB 1 GTCACTTTGGGAGCCGGGCATCGCTGTCGCGCCAGGAGCCTGCCAGGAGGC
QY 289 GCAGAGGAGGACCCAGGACCGGTGCGAACTGAATCCCGACAGAGAAAGCCAGG;
DB 61 GCAGAGGAGGACCCAGGACCGGTGCGAACTGAATCCCGACAGAGAAAGCCAGG;
QY 349 GCGCCTTTCTGAAACCGACTAGTTCGGCTCGCAGAGTGACCTAAAGGCGGGA
DB 121 GCGCCTTTCTGAAACCGACTAGTTCGGCTCGCAGAGTGACCTAAAGGCGGGA
QY 409 CGGGCTCGAAGAGCGATCGAGCCCATTTATGAAGTTTCATCCACGACCTGGACAGG
DB 181 CGGGCTCGAAGAGCGATCGAGCCCATTTATGAAGTTTCATCCACGACCTGGACAGG
QY 469 GCGCAGGCGAGGTGGACGGGACAGTGTAGTGGCTGGGAGGAGCCAGAAATCAACA
DB 241 GCGCAGGCGAGGTGGACGGGACAGTGTAGTGGCTGGGAGGAGCCAGAAATCAACA
QY 529 AGCCCTCTGCGCTACAACCGCCAGATCGGGAGTTTATAGTCACCCGGGCTGGGCT
DB 301 AGCCCTCTGCGCTACTACCGCCAGATCGGGAGTTTATAGTCACCCGGGCTGGGCT
QY 589 TACCTGTACTGTCAAGGTGCATTTTGTATGAGGGGAGGCTGTCTACCTGAAGCTGG
DB 361 TACCTGTACTGTCAAGGTGCATTTTGTATGAGGGGAGGCTGTCTACCTGAAGCTGG
QY 649 CTGTGTGATGTGTGTGCGCCCTGGCTGCGCTGGAGGAAATTCAGCCACTGGCG
DB 421 CTGTGTGATGTGTGTGCGCCCTGGCTGCGCTGGAGGAAATTCAGCCACTGGCG
QY 709 TCCCTCGGGGCCCCAGCTCCGCTCTGCGCAGGTGTCTGGGCTGTGGGCTGGGCT
DB 481 TCCCTCGGGGCCCCAGCTCCGCTCTGCGCAGGTGTCTGGGCTGTGGGCTGGGCT
QY 769 TCCCTCGGGGCCCCAGCTCCGCTCCCTGGGCCCCATCTCAAGGCTGGCCCCCTTCT
DB 541 TCCCTCGGGGCCCCAGCTCCGCTCCCTGGGCCCCATCTCAAGGCTGGCCCCCTTCT
QY 829 TACTTCGAGCTTTCCAGGTTCACTGAGGGGCCCTGTGTCTCCCCACAGTCTCCCP
DB 601 TACTTCGAGCTTTCCAGGTTCACTGAGGGGCCCTGTGTCTCCCCACAGTCTCCCP
QY 889 GCCGGCTCCCTCTGACAGCTCTCTGCGGCGACCCGGTCCCTCTGCCCCACCTCAG
DB 661 GCCGGCTCCCTCTGACAGCTCTCTGCGGCGACCCGGTCCCTCTGCCCCACCTCAG

06:25:17 2004

us-09-245-198a-3.rng

GTCTCAGACCTGCCCTCCCTCTAGAGCGTCTGGGCTGTTCACGTGTTTCCA 100B
 GTCTCAGACCTGCCCTCCCTCTAGAGCGTCTGGGCTGTTCACGTGTTTCCA 780
 ACATAATACAGTATTCCTCACTTATCTTCAACTCCCCACCGCCCACT 1063
 ACATAATACAGTATTCCTCACTTATCTTCAACTCCCCACCGCCCACT 835

dard; cDNA; 1239 BP.

(first entry)

AK coding sequence.

EAK; TNF relatedness and weak ability to induce cell death; necrosis factor; TWEAK; fibrosis; cardiac disease; lung disease; kidney disease; skin disease; le disease; adipose tissue disease; nal tract disease; pancreatic disease; organ disease; neural disease; cartilage disease; connective tissue disease; cellular death; hepatotropic; gastrointestinal; osteopathic; gene; ss.

```
Location/Qualifiers
1. .750
/*tag= a
/product= "FL-TWEAK"
```

32.

2003WO-US011350.

2002US-0371611P.

J INC.

ubowski A, Zheng T, Hahn K:

:56/78.

27

ic, cartilage or neural tissue condition in a subject nistering to the subject a TWEAK agonist or antagonist.

) ID NO 2; 120pp; English.

sequence is the coding sequence for murine transmembrane FL-attenuates and weak ability to induce cell death, where TNF (toxic factor). TWEAK is a member of the TNF family. TWEAK antagonists are useful for treating a TWEAK-related condition, such as fibrosis; cardiac disease; liver disease; lung disease; skin disease; skeletal muscle disease; adipose tissue disease; gastrointestinal tract disease; pancreatic disease; reproductive system disease; cartilage disease; bone disease; nerve disease; cellular death; and a pathological condition involving a TWEAK receptor.

BP; 249 A; 386 C; 331 G; 273 T; 0 U; 0 Other;

49.8%; Score 683.4; DB 9; Length 1239;

77.08; Pred. No. 9.7e-136;

conservative 0; Mismatches 221; Indels 70; Gaps 9;

QY	106	ATGGCGCCGCTTCGAGCCACAGAGCGAGGGGGGCGCCCGGGGGGAGCCCGGGCACCG
DB	1	ATGGCGCCGCTTCGAGCCACAGAGCGAGGGGGCGCGGGGGAGCGCGGCACCG
QY	166	CTGTGCCGCTCGCCTGGGCTTGGGCTTGCGCTGGCTTGCCTCGCTCGGCTCTCGCT
DB	61	CTGGCCCCGCTGGTGTGAGCCTTGGGCTTGCGCTGGCTTGCCTTGGCTCTCGCT
QY	226	GTTGTCAGTTTTGGGAGCGCGGCATCGCTGTCGCGCCAGGAGCCTGCCACGAGG
DB	121	GTTGTCAGCCTTGGGAGCTGGCAACGCTGTCTGCCAGGAGCCTTCTACGAGG
QY	286	GTCGACAGGAGGACCGAGACCCTGTCGGAACGTAAATCCCACAGAGAAGAACCT
DB	181	ACAGCAGAGGACCGCGGAGCCCTGAACTGAATCCCCCAGACAGAGGAAAGCC
QY	346	CCTCGCCTTTCCTGAACCGACTAGTTTGGGCTTCGAGAGTGACCTTAAGGGCC
DB	241	GTGGTA CTTTCTTTGGAAACAATAAGTCCGCGCTCGAAGAA GTCTCTTAAGGCC
QY	406	ACCGGCTCGAAGAGCGATCGCAGGCCATTATGAACTTCATCCACGACCTGGACA
DB	301	GC CGGCTCGCCGAGCTATTGCAGCCATTATGAGGTT CATCTCGGCGAGGACA
QY	466	GGAGCGAGGACGCTGTGSACGGSACAGTAGTGGCTGGGAGGAAGCCAGAACTCAA
DB	361	GGAGCA AAGCAGGTGGATGGACAGTAGTGGCTGGGAGAGACCAAAATCAA
QY	526	TCCAGCCCTCTGGCTACAACCGCCAGATCGGGAGTTTATAGTACCCCGGCTGG
DB	421	TCCAGCCCTCTGGCTACGACCGCCAGATTGGGGAATTTACAGTCATCAGGGCTGG
QY	586	TACTACTGTACTGTCAAGTGCATTTTGATGAGGGAAGGCTCTCTACCTGAAGCT
DB	481	TACTACTGTACTGTCAAGTGCATTTTGATGAGGGAAGGCTGTCTACCTGAAGCT
QY	646	TTGCTGTGTGATGGTGTGTGGCCCTCGCTGCTGGAGGAA TTTCTCAGGCCACTGC
DB	541	TTGCTGTGAACGGTGTGCTGGCCCTGGCTGCTGGAAGAA TTCTCAGCCA CAGC
QY	706	AGTTCCCTGGGCCCCAGCTCGCGCTCTGCGCTGCTGGAGGAGTTGTGGCCCTGGCG
DB	601	AGCTCTCTGGGCCCCAGCTCCGTTTGTGCCAGTGTCTGGGCTGTTCGGCTGGCG
QY	766	GGGTCTCTCCCTGGGATCGSACCCCTCCCTGGGCCCCATCTCAAGGCTGCCCGCTT
DB	661	GGGTCTCTCCCTCGGATCGSACCCCTCCCTGGGCTCATCTTAAAGGCTGCCCGCTT
QY	826	ACCTACTTCGGACTCTTCAGGTTTCACTGAGGGGCCCTGTCTCTCCCA CAGTCGTC
DB	721	ACCTACTTTGGACTCTTCAAGTTTCACTGAGGGGCCCTTGTCTCTCCAGATTCCTTA
QY	886	GCTGCCGGCTCC-----CCTCGACAGCTCTTGGGCA CCGGTCCTCTGGCCCCA
DB	781	TTCCCTTGGCTCCAGGAGCATCACACCTCCCTACCCCA CCCCCACTCTCTCCACC
QY	941	CAGCCGCTCTTTGCTCCAGACCTGCCCTCCCTCTAGAGGCTGCGCTTGGGCTGTCT
DB	841	C-GCTGCTCTTGGTCCATGCTGTCTCTCC--TAAAGGCGAGCAGAGCTTGTCT
QY	1001	GTTTTCCATCCCAATAAATACAGTATCCCACTCTTATCTTACAACTCCCCCA CC
DB	898	GTTTTCCATCC-----ACAGACGTATCTTGTCTTCTTAACATCCCATCCCA CT
QY	1061	ACTCTCCACTCACTAGTCTCCCAATTCCTTGACCTTTT GAGGCCCCCCAGTGTCT
DB	951	ACTATCCACTCACTAGCTCCAAAAGCCCTAC-----TTATCCCTCT
QY	1121	CCCCCTGGCCACAGACCCCGAGGCAATGTGTTTCACTGTACTCTGTGGGCAAGAT
DB	996	CCCCCA CCACTCACCGGACCACTGTTTATTGACTTTGTGCAC-----
Y	1181	TCCAGAAAGCCCCACTTTCAGGCATTAAGAGGGGCTTGGACCTCGCGGCGAGGACCA

-----CAGGCACTGAGATGGGCTGGACCTGGTGGCAGGAAGCCAGAGA 1082
TGGCCCTAGGCGAGGAGTTCCCAATGTGAGGGGCGAGA-AACAAGACAAGCTCCTC 1299
TGGGACTAGGCGAGGAGTTCCCACTGTGAGGGGGAAGAGCTGGGGACAAGCTCCTC 1142
TGGAATTCCTGTGGATTTTAAACACAGATATTATTTTATTTATTTATTTGACAAA 1359
GGA-----TCCCTGTGGATTTTGAAG--AGATACTATTTTATTTATTTATTTGACAAA 1196
TTGA 1366
TTAA 1203

standard; DNA; 898 BP.

(first entry)

vector pDC409-LZ-TWEAK fusion protein-encoding DNA.

cellular domain; tumour necrosis factor; TNF; angiogenesis;
sclerolisation; diabetic retinopathy; neovascular glaucoma;
na; retinopathy of prematurity; retrolental fibroplasia;
itis; macular degeneration; arthritis; rheumatism; ds;
neovascularisation; psoriasis; metastatic condition;
mour; sarcoma; carcinoma; benign tumour; haemophilic joint;
condition; myocardial angiogenesis; ischaemia; human;
vascular adhesion; telangiectasia; wound granulation;
ic plaque neovascularisation; coronary atherosclerosis;
therosclerosis; pDC409-LZ-TWEAK; TWEAK receptor; TWEAKR;
n.

Location/Qualifiers

52..873

/tag= a

/product= "Fusion protein comprising a growth hormone
leader, a leucine zipper multimerisation domain, and
human TWEAK extracellular domain"

12.

2000WO-US034755.

99US-0172878P.

2000US-0203347P.

TEX CORP.

975/44.
-499.

angiogenesis in a mammal for treating diseases mediated by
e.g. solid tumors and vascular deficiencies of cardiac or
issue, by administering antagonist or agonist of TWEAK

ge 39-40; 46pp; English.

represents a DNA from the expression vector pDC409-LZ-TWEAK,
a fusion protein comprising a growth hormone leader, a

CC leucine zipper multimerisation domain, and the extracellular dom
CC human TWEAK. The fusion protein was used in the isolation of hum
CC receptor (TWEAKR)-expressing clones from a COS cell human CDNA l
CC The TWEAK protein is a member of the tumour necrosis factor (TNF
CC and induces angiogenesis. TWEAKR may therefore be used to screen
CC develop TWEAKR agonists and antagonists for the modulation of
CC angiogenesis, to be used in the treatment and diagnosis of human
CC The disorders mediated by angiogenesis include ocular disorders
CC characterised by ocular neovascularisation such as diabetic reti
CC neovascular glaucoma, retinoblastoma, retinopathy of prematurity
CC retrolental fibroplasia, rubeosis, uveitis, macular degeneration
CC corneal graft neovascularisation, and inflammatory diseases such
CC arthritis, rheumatism and psoriasis. Other treatable diseases in
CC malignant and metastatic conditions such as sarcomas and carcino
CC benign tumours and preneoplastic conditions, myocardial angione
CC haemophilic joints, scleroderma, vascular adhesions, atheroscler
CC plaque neovascularisation, telangiectasia, wound granulation, co
CC atherosclerosis, peripheral atherosclerosis and ischaemia
XX
XX
SQ Sequence 898 BP; 187 A; 266 C; 267 G; 178 T; 0 U; 0 Other;

Query Match 45.8%; Score 629.2; DB 4; Length 898;
Best Local Similarity 99.5%; Pred. No. 3.1e-124;
Matches 631; Conservative 0; Mismatches 3; Indels 0;

QY 232 AGTTTGGGAGCGGGCATCGCTGTCGCGCCAGGAGCCTGCCAGGAGAGCTGG
Db 250 AGTTTGGGAGCGGGCATCGCTGTCGCGCCAGGAGCCTGCCAGGAGAGCTGG
QY 292 GAGGAGGACAGGACCCGTCGGAATGAATCCCGAGAGAGAAAGCCAGGATC
Db 310 GAGGAGGACAGGACCCGTCGGAATGAATCCCGAGAGAGAAAGCCAGGATC
QY 352 CTTTCTCCGACCGGACTAGTTCCGCTCGCAGAGTGCACCTAAAGGCCGGA
Db 370 CTTTCTCCGACCGGACTAGTTCCGCTCGCAGAGTGCACCTAAAGGCCGGA
QY 412 GCTCGAAGAGCGGATCGCAGCCCATTTATCAAGTTTCATCCAGACCTGGACAGGACG
Db 430 GCTCGAAGAGCGGATCGCAGCCCATTTATCAAGTTTCATCCAGACCTGGACAGGACG
QY 472 CAGGAGGTGTGGACGGACAGTGTGCTGGGAGGAGCCAGAGATCAACAGCT
Db 490 CAGGAGGTGTGGACGGACAGTGTGCTGGGAGGAGCCAGAGATCAACAGCT
QY 532 CCTCTGCGCTACACCGCCAGATCGGCGAGTTTATAGTCACCCGGGCTGGGCTCT
Db 550 CCTCTGCGCTACACCGCCAGATCGGCGAGTTTATAGTCACCCGGGCTGGGCTCT
QY 592 CTGTACTGTCAAGTGTCACTTTTGTATGAGGGGAGGCTGTCTACCTGAAGCTGGACT
Db 610 CTGTACTGTCAAGTGTCACTTTTGTATGAGGGGAGGCTGTCTACCTGAAGCTGGACT
QY 652 GTGATGTGTGTGGCCCTCGGCTGTGCGAGGTGTGGGCTGTGGCCCTGCGGCCAG
Db 670 GTGATGTGTGTGGCCCTCGGCTGTGCGAGGTGTGGGCTGTGGCCCTGCGGCCAG
QY 712 CTGCGGCCCGCCAGCTCCGCTCTGCGAGGTGTGGGCTGTGGCCCTGCGGCCAG
Db 730 CTGCGGCCCGCCAGCTCCGCTCTGCGAGGTGTGGGCTGTGGCCCTGCGGCCAG
QY 772 TCCCTGGCGGATCCCGCACCCCTCCCTGGGCCCATCTCAAGGCTGCCCTTCTCTCA
Db 790 TCCCTGGCGGATCCCGCACCCCTCCCTGGGCCCATCTCAAGGCTGCCCTTCTCTCA
QY 832 TTGCGATCTTTCAGGTTACTGTAGGGGCCCTGG 865
Db 850 TTCGGAATCTTCCAGGTTCACTGAGGCGCGCGG 883

RESULT 14

AAV18599

ID AAV18599 standard; cDNA; 1168 BP.

CTGGGCAACGCTGTCTGCCAGGAGCGTTCTCAGGAGAGCTCACAGCGACCGC 121

[illegible]

ndard; DNA; 701 BP.

(first entry)

DNA.

sis factor receptor; signal transducer molecule; TNF; APO4;
 1 abnormality; gestational abnormality; prostate cancer;
 APO4; TNRL-1; TNRL-3; diagnosis; treatment; therapy; disease;
 domain; immunogen; antibody preparation; breast carcinoma;
 cause; ss.

Location/Qualifiers
 1. .636
 /*tag= a
 /product= "TNRL3"

98WO-US018393.
 97US-00924634.
 WASHINGTON.

3191/17.
 1591.

rosis Factor family receptor polypeptides and ligands -
 agnosis and treatment of prostate cancer and developmental
 al abnormalities.

Fig 13B; 156pp; English.

n describes isolated Tumor Necrosis Factor (TNF) family
 peptides: APO4, APO6, APO8 and APO9 or their active
 id isolated TNF related ligands 1 and 3 (TNRL1 and TNRL3) or
 fragments. APO4 is useful for diagnosing prostate cancer by
 levels of APO4 in an individual. Prostate cancer can also be
 APO4 selective binding agents linked to a therapeutic
 polypeptides are also useful for identifying selective
 s, useful in diagnosis/treatment of disease by binding of
 polypeptide/active fragment which is extracellular, or
 the cell surface. The binding is preferably performed in
 polypeptides/ active fragments are also useful for screening
 and antagonists by binding and observing the change in APO4
 active pharmacological agents useful in diagnosis or
 disease are also identified using APO4 polypeptides/active
 APO4 signal transducer molecules that specifically interact
 asmic domain of APO4 and detecting a change in level of APO4
 method is performed in vivo or in vitro. APO polypeptides
 l as immunogens for preparing antibodies. APO4 is also
 agnosis/treatment of developmental or gestational
 . APO8 was transfected to human breast carcinoma cell line
 duced apoptosis

BP; 139 A; 210 C; 203 G; 149 T; 0 U; 0 Other;
 37.8%; Score 519.2; DB 2; Length 701;
 arity 87.3%; Pred. No. 7.3e-101;
 onservative 0; Mismatches 83; Indels 0; Gaps 0;

220 CTGGCGTGGTCTAGTTTGGGAGCCGGGAGTCGCTGTCCGCCAGGAGCGCTGCC
 1 CTGGTGGTGGTCTAGCTTGGGAGCTGGGCAACGCTGTCTGCCAGGAGCGCTTCTC
 280 GAGCTGGTGGCAGAGGAGGACCCGTCGGAACCTGAATCCCGCAGACAGAAAG
 61 GAGCTGACAGCAGAGGAGCCCGGGAGCCCGCTGAATGAATCCCGCAGACAGAGG
 340 CAGGATCCCTGGCCCTTCTGTAACCGACTAGTTCGGCCTCGCAGAAAGTGACCTA
 121 CAGGATGGTACCTTCTTGGAACTAGTTCGGCCTCGAAGAAAGTGCTCTA
 400 CGGAAACACGGGCTCGAAGAGCGATCGAGCCATTATGAAGTTTCATCCAGAC
 181 CGGAAGCGGGCCTCGCCGAGCTATTGAGCCCATTAAGGTTTCATCTCTGGG
 460 CAGGACGGAGCGCAGCGAGGTGTGGAACGGGACAGTGAGTGGCTGGGAGGAGCCA
 241 CAGGATGGAGCACAGCAGGTGTGGATGGGACAGTGAGTGGCTGGGAGGAGCCA
 520 AACAGCTCCAGCCCTCTGGCTACACCGCCAGATCGGGAGGTTTATAGTACCC
 301 AACAGCTCCAGCCCTCTGGCTACACCGCCAGATTTGGGAAATTTACAGTCA
 580 GGGCTCTACTACTGTACTGTCTGAGTGCATTTGATGAGGGGAGGCTCTCTACC
 361 GGGCTCTACTACTGTACTGTCTGAGTGCATTTGATGAGGGGAGGCTCTCTACC
 640 CTGGAATTGCTGGTGGATGGTGTCTGGCCCTCGCTGCTGGAGGAATTTCTCAG
 421 CTGGAATTGCTGGTGGATGGTGTCTGGCCCTCGCTGCTGGAGGAATTTCTCAG
 700 GGGGCCAGTTCCCTCGGGCCCCCAGCTCCGCTCTGCGAGGTGTCTGGGCTGTGG
 481 GAGCAAGCTCTCTCTGGGCCCCCAGCTCCGCTTTGTGCCAGGTGTCTGGGCTGTGG
 760 CGGCCAGGGTCTCTCTCGGATCCGACCCCTCCGCTGGGCCCCATCTCAAGGCTG
 541 CGGCCAGGGTCTCTCTCGGATCCGACCCCTCCGCTGGGCTCATCTTAAGGCTG
 820 TTCCTCAGCTTCTCGGATCTTCCAGGTTCACTAGGGGGCCCTGGTCTCCC 87
 601 TTCCTAAGCTTCTTGGACTCTTTCAAGTTCAGTTCAGTGGGGGCTTGTCTCTCCC 65

Search completed: April 7, 2004, 21:32:21
 Job time : 528.669 secs

3	0	1364	9	US-09-822-849A-19	Sequence 19, Appl
3	0	1363	14	US-10-210-951-3	Sequence 3, Appl
3	0	1353	14	US-10-211-894-3	Sequence 3, Appl
3	0	1306	12	US-10-202-062-23	Sequence 23, Appl
3	0	1306	14	US-10-272-411-16	Sequence 16, Appl
3	0	1306	14	US-10-218-547-23	Sequence 23, Appl
3	0	1306	14	US-10-272-358A-16	Sequence 26, Appl
3	0	898	9	US-10-310-793-29	Sequence 29, Appl
3	0	898	9	US-09-742-454A-1	Sequence 1, Appl
3	0	898	9	US-09-883-777-1	Sequence 1, Appl
3	0	493	10	US-09-918-995-2125	Sequence 21225, A
3	0	493	9	US-09-960-352-2197	Sequence 2197, Ap
3	0	7	213	US-10-085-783A-55176	Sequence 55176, A
3	0	7	213	US-10-242-535A-55176	Sequence 55176, A
3	0	264	9	US-09-983-965-2183	Sequence 2183, Ap

ALIGNMENTS

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RESULT 1
US-09-822-849A-19
/ Sequence 19, Application US/09822849A
/ Patent No. US20020045170A1
/ GENERAL INFORMATION:
/ APPLICANT: Wong, Gordon G.
/ APPLICANT: Clark, Hilary
/ APPLICANT: Fechtel, Kim
/ APPLICANT: Agostino, Michael J.
/ APPLICANT: Howes, Steven H.
/ APPLICANT: Resnick, Richard J.
/ APPLICANT: Gulkota, Kamalakara
/ APPLICANT: Graham, James R.
/ APPLICANT: Genetics Institute, Inc.
/ TITLE OF INVENTION: POLYNUCLEOTIDES ENCODING NOVEL SECRETED PROTEIN
/ FILE REFERENCE: GIN 6403
/ CURRENT APPLICATION NUMBER: US/09/822,849A
/ CURRENT FILING DATE: 2001-09-04
/ PRIOR APPLICATION NUMBER: 60/195,582
/ PRIOR FILING DATE: 2000-04-06
/ NUMBER OF SEQ ID NOS: 598
/ SOFTWARE: PatentIn Ver. 2.0
/ SEQ ID NO 19
/ LENGTH: 1364
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-822-849A-19

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Query Match      98.0%; Score 1345.8; DB 9; Length 1364;
Best Local Similarity 99.9%; Pred. No. 0;
Matches 1347; Conservative 2; Indels 0; G:
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5 TC CGC CGC CGC GGT CCG CCG TCC CGC GAT CCG TCG GTC CCG CGG ATCG GGG GGG CGGT
85 CAG GCA CAG CCG CCG CCG CCG CCG TCG GAC CAG CAG GGG GGG GGG GCG

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[illegible]

RESULT 2
US-10-210-951-3
; Sequence 3, Application US/10210951
; Publication No. US20030170228A1
; GENERAL INFORMATION:
; APPLICANT: Ashkenazi, Avi J.
; APPLICANT: Goddard, Audrey
; APPLICANT: Godowski, Paul J.
; APPLICANT: Gurney, Austin L.
; APPLICANT: Hillan, Kenneth J.
; APPLICANT: Marsters, Scot A.
; APPLICANT: Pan, James
; APPLICANT: Pitti, Robert M.
; APPLICANT: Roy, Margaret Ann
; APPLICANT: Smith, Victoria
; APPLICANT: Stone, Donna M.
; APPLICANT: Watanabe, Colin K.
; APPLICANT: Wood, William I.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE TREATMENT OF
; FILE REFERENCE: P2931R1C1

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; CURRENT APPLICATION NUMBER: US/10/210,951
; CURRENT FILING DATE: 2002-08-02
; PRIOR APPLICATION NUMBER: 60/014699
; PRIOR FILING DATE: 1996-04-01
; PRIOR APPLICATION NUMBER: 60/026943
; PRIOR FILING DATE: 1996-09-23
; PRIOR APPLICATION NUMBER: 60/059121
; PRIOR FILING DATE: 1997-07-17
; PRIOR APPLICATION NUMBER: 60/059352
; PRIOR FILING DATE: 1997-09-19
; PRIOR APPLICATION NUMBER: 60/062037
; PRIOR FILING DATE: 1997-10-10
; PRIOR APPLICATION NUMBER: 60/063755
; PRIOR FILING DATE: 1997-10-17
; PRIOR APPLICATION NUMBER: 60/063045
; PRIOR FILING DATE: 1997-10-24
; PRIOR APPLICATION NUMBER: 60/063046
; PRIOR FILING DATE: 1997-10-24
; PRIOR APPLICATION NUMBER: 60/066511
; PRIOR FILING DATE: 1997-11-24
; PRIOR APPLICATION NUMBER: 60/066772
; PRIOR FILING DATE: 1997-11-24
; Remaining Prior Application data removed - See File Wrapper or PAL
; NUMBER OF SEQ ID NOS: 258
; SEQ ID NO 3
; LENGTH: 1353
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-210-951-3

Query Match          96.2%; Score 1320.2; DB 14; Length 1353;
Best Local Similarity 99.8%; Pred.No. 0;
Matches 1322; Conservative 0; Mismatches 3; Indels 0;

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| | | | |
Ddb     1    CGATCCCTCGGGTCCCGGATGGGGGGGGGTAGGCAGGCACACCCCCCGCCC
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Qv      109  GCGCSCCGTCGAGCACAGAGGCGAGGGGGGGCCCGGGGGAGCTGGGACACCGCCT

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3CCGTCGGAGCCHAGAGCGAGGGGGCGCCGGGGGGAGCCGGGCAACCGCCTCTCGT 168

3CCCGTCGGAGCCHAGAGCGAGGGGGCGCCGGGGGGAGCCGGGCAACCGCCTCTCGT 120

3CGCTCGCGCTGGGCGCTGGGCGCTGGCCCTGGCTCGCCTCGGCTCTCTGCTGGCCGCTG 228

3CGCTCGCGCTGGGCGCTGGGCGCTGGCCCTGGCTCGGCTCTCTGCTGGCCGCTG 180

AGTTTGGGAGCCGGGGCATCGCTGTCCGCCCCAGGAGCGCTGCCCAAGGAGGAGCTGGT 288

AGTTTGGGAGCCGGGGCATCGCTGTCCGCCCCAGGAGCGCTGCCCAAGGAGGAGCTGGT 240

3AGGAGACAGGACCCGTCGGAACTGAATCCCCAGACAGAAAGAACGCCAGGATCCCT 348

3AGGAGACAGGACCCGTCGGAACTGAATCCCCAGACAGAAAGAACGCCAGGATCCCT 300

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CCTTTCTGGAACCGACTAGTTTCGGCTCGCAGAGAGTGCACTTAAGGCCCGGAAAAACA 360

3CTCGAAGACGATCGCAGGCCATTATGAAGTTTCATCCAGACCTGCACAGGACCGGA 468

3CTCGAAGACGATCGCAGGCCATTATGAAGTTTCATCCAGACCTGCACAGGACCGGA 420

3AGCAGGTGTGACGGGACAGTAGTGGCTGGGAGGAGCCAGAACTCAACAGCTCC 528

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3TGGGGCCCGAGCTCGCCCTCTGCCAGTGTCTGGGCTGTGGGCCCTGGCCGACGG 720

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3CCCTCGGATCGCAGCCCTCCCTGGGCCCATCTCAAGGCTGCCCTCTCTCAACC 780

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3CACTTCAGGCACCTAAGAGGGGCTGGACCTGGGGGCGAGGAAGCCAAAGAGACTGGGC 1248

[illegible]

Query Match	93.6%	Score 1285;	DB 12;	Length 1306;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 1285;	Conservative 0;	Mismatches 0;	Indels 0;	
QY	89	CACAGCCCCCGCCCCCATGSCGCCCGCTCGAGCCNAGAGCGCGAGGGGGCGCGCG		
Db				
	1	CACAGCCCCCGCCCCCATGSCGCCCGCTCGAGCCNAGAGCGCGAGGGGGCGCGCG		
QY	149	AGCGGGGACACCGCCTGCTGTGTCCCGCTCGCGCTGGGSCCTGGGSCCTGGCGCTGGCG		
Db				
	61	AGCGGGGACACCGCCTGCTGTGTCCCGCTCGCGCTGGGSCCTGGGSCCTGGCGCTGGCG		
QY	209	TGCGGCTCTCTGCTGGCGGTGGTCAGTTTGGGGAGCGGGGCATCGCTGTCCGGCCCA		
Db				
	121	TGCGGCTCTCTGCTGGCGGTGGTCAGTTTGGGGAGCGGGGCATCGCTGTCCGGCCCA		
QY	269	CTGCCCCAGGAGAGCTGTGTGGCAGAGAGGACACAGGACCCGTCGGAACCTGAATCC		
Db				
	181	CTGCCCCAGGAGAGCTGTGTGGCAGAGAGGACACAGGACCCGTCGGAACCTGAATCC		
QY	329	CAGAAAGAACCGAGATCTCTGGCGCTTTCTCTGAAACCGACTAGTTTGGCGCTCGCAG		
Db				
	241	CAGAAAGAACCGAGATCTCTGGCGCTTTCTCTGAAACCGACTAGTTTGGCGCTCGCAG		
QY	389	CACCTAAAGGCGGAAACACGCGCTCGAAGAGCGATCGCAGCCCATTTATGAAGT		
Db				
	301	CACCTAAAGGCGGAAACACGCGCTCGAAGAGCGATCGCAGCCCATTTATGAAGT		
QY	449	CACGACCTTGGACAGGACGAGCGCAGCGAGGTTGGACGGGACAGTGAATGGCTG		
Db				
	361	CACGACCTTGGACAGGACGAGCGCAGCGAGGTTGGACGGGACAGTGAATGGCTG		
QY	509	AGGCCAGAGATCAACAGCTCCAGCGCCTCTCGGCTTACAAACCCAGATCGGGGAGTT		
Db				
	421	AGGCCAGAGATCAACAGCTCCAGCGCCTCTCGGCTTACAAACCCAGATCGGGGAGTT		
QY	569	TCACCCGGGCTGGGCTCTACTACTGTCTCAGGTGACCTTTGATGAGGGGAA		

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TGGCCCTCGGGCCAGGGTCTCTCCCTCGCGATCCGACACCTCCCTCGGGCCCATCTCA	720
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CTGCCCCCTTCTCTACCTACTTCTGGACTCTTCACAGTTCACTGAGGGCCCTGGTCT	780
CAAGTCGTCTCCAGAGCTGCCGGCTCCCTCCGACAGCTCTCTGGGACCCCGTCCCTT	928
CAAGTCGTCTCCAGAGCTGCCGGCTCCCTCCGACAGCTCTCTGGGACCCCGTCCCTT	840
CCCCACCTCAGCGCTCTTTGTCTCCAGACTGCCCTCCCTCTAGAGGCTCCCTGG	988
CCCCACCTCAGCGCTCTTTGTCTCCAGACTGCCCTCCCTCTAGAGGCTCCCTGG	900
TGTTCCAGTGTTTTCCATCCACACATAAATACAGTATCCCACTCTTATCTTACAAC	1048
TGTTCCAGTGTTTTCCATCCACACATAAATACAGTATCCCACTCTTATCTTACAAC	960
CCACGGCCCACTCTCCACCTCACTAGTCTCCCAATCCCTGACCCCTTTGAGGCCCCCA	1108
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AGCCAAAGAGACTGGGCTTAGCCAGAGTTCCTCCAAATGTGAGGGCGAGAAACAAG	1200
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GTGACAAATGTTGATTAATGG	1373
GTGACAAATGTTGATTAATGG	1285

Publication US/10218547
US20030100074A1
[TITON:
an Genome Sciences, Inc.
[TITON: Methods And Compositions For Treating Metabolic Bone Diseases Rel
[TITON: Human Endokine Alpha
: PF561
[TITON NUMBER: US/10/218,547
DATE: 2002-08-15
[TITON NUMBER: 60/312,542
ATE: 2001-08-15
[TITON NUMBER: 60/330,761
ATE: 2001-10-30
D NOS: 57
it in version 3.1

10

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93.6%; Score 1285; DB 14; Length 1306;
arity 100.0%; Pred. No. 0;
'conservative 0; Mismatches 0; Indels 0;

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QY	89	CACAGCCCCCGCCCGCCGATGGCGCCCGTGGAGCCAGAGCGAGCGGGGCGCGC
DB	1	CACAGCCCCCGCCCGCCCATGGCGCCCGTGGAGCCAGAGCGAGCGGGGCGCGC
QY	149	AGCGGGACACGCCCTGCTGCTCCCGCTCGCGCTGGGCTCGGGCCTCGCGCTGG
DB	61	AGCGGGACACGCCCTGCTGCTCCCGCTCGCGCTGGGCTCGGGCCTCGCGCTGG
QY	209	TCGGCTCTCTGTGGGCCGTGSGTCACTTTTGGGAGCCGGGCATCGCTTCGGCCCG
DB	121	TCGGCTCTCTGTGGGCCGTGSGTCACTTTTGGGAGCCGGGCATCGCTTCGGCCCG
QY	269	CTGCCACAGAGGAGCTGTGTGGCAGAGGAGGACAGAGCCCGTCGGAACTGAATCC
DB	181	CTGCCACAGAGGAGCTGTGTGGCAGAGGAGGACAGAGCCCGTCGGAACTGAATCC
QY	329	CAGAAGAAAGCCAGGATCCGTGGCCTTTCCTGAAACCGACTAGTTCGGCCTCGCAC
DB	241	CAGAAGAAAGCCAGGATCCGTGGCCTTTCCTGAAACCGACTAGTTCGGCCTCGCAC
QY	389	CACCTAAAGGCCGGA AAA CA CGGGCTCGAAAGAGCGATCGCAGGCCCATATGAAGT
DB	301	CACCTAAAGGCCGGA AAA CA CGGGCTCGAAAGAGCGATCGCAGGCCCATATGAAGT
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DB	361	CAGGACCTCGA CAGGACGAGCGCAGGCGAGGTGTGGA CGGGA CAGTGA GTGGCTC
QY	509	AAGCCAGAAATCAACAGCTCCAGCCCTCGCGCTACAAACCGCAGATCGGGGAGTT
DB	421	AAGCCAGAAATCAACAGCTCCAGCCCTCGCGCTACAAACCGCAGATCGGGGAGTT
QY	569	TCACCCGGGCTGGGCTCTACTACCTGTACTGTCAAGTGCACCTTTGATGAGGGGAA
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QY	749	TGTTGGCCCTGCGGCCAGGGTCTCTCCCTGCGGATCGGCACCCCTCCCTCGGGCCCA
DB	661	TGTTGGCCCTGCGGCCAGGGTCTCTCCCTGCGGATCGGCACCCCTCCCTCGGGCCCA
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QY	869	CCCCACAGTCTGCCAGGCTGCGGCTCCCTCGACAGCTCTCTGCGGCACCCCGGT
DB	781	CCCCACAGTCTGCCAGGCTGCGGCTCCCTCGACAGCTCTCTGCGGCACCCCGGT
QY	929	CTGCCACACCTCTAGCCGCTCTTTGTCTCCAGACTCCGCCCTCCCTCTAGAGGCTG
DB	841	CTGCCACACCTCTAGCCGCTCTTTGTCTCCAGACTCCGCCCTCCCTCTAGAGGCTG
QY	989	GCGCTTTCAGGTGTTTCCATCCCAATAAATACAGTATTCCTACTCTTATCTTAT
DB	901	GCGCTTTCAGGTGTTTCCATCCCAATAAATACAGTATTCCTACTCTTATCTTAT
QY	1049	CCCCACCGGCCACTCTCCACCTCACTAGCTGCCCAATCCCTGACCCCTTGAGGCT
DB	961	CCCCACCGGCCACTCTCTCACCTCACTAGCTGCCCAATCCCTGACCCCTTGAGGCT
QY	1109	GTGATCTCGACTCCCCCTGGCCACAGAGCCCGCAGGCAATGTGTTCCTACTGCT
DB	1021	GTGATCTCGACTCCCCCTGGCCACAGAGCCCGCAGGCAATGTGTTCCTACTGCT
QY	1169	GGCAAGGATGGGTCAGAAAGACCCCACTTCAGGCACTAAGAGGGGCTCGGACCTGGC

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: TITLE OF INVENTION: Methods And Compositions For Treating Inflammato
:
: TITLE OF INVENTION: Relating To Human Tumor Necrosis Factor-Gamma I
: FILE REFERENCE: PF573
:
: CURRENT APPLICATION NUMBER: US/10/310,793

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Db	961	CCCCACGGCCACTCTCCACTCACTAGCTCCCAATCCCTGACCTTTGAGGC
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Db	1021	GTGATCTGACTCCCTCCCTGTGCCACAGACCCCGAGGCAATTGTTCACGTACT
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Db	1081	GCGAAGGATGGTCCAGAGAAGACCCCACTTCAGGCACTAAGAGGGGCTGCACTGG
Qy	1229	GGAAAGCAAGAGACTGGGCTTAGGCCAGAGATCCCAATGTGAGGGGCGAGAA
Db	1141	GGAGCCAAAGAGACTGGGCTTAGGCCAGAGATCCCAATGTGAGGGGCGAGAA
Qy	1289	ACAAGCTCCTCCTTGAGAAATTCCTGTGATTTTTTAAACAGATATTATTTTAA
Db	1201	ACAAGCTCCTCCTTGAGAAATTCCTGTGATTTTTTAAACAGATATTATTTTAA
Qy	1349	ATTGTGACAAAATGTTGATAATGG 1373
Db	1261	ATTGTGACAAAATGTTGATAATGG 1285

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RESULT 9
US-09/742-454A-1
; Sequence 1, Application US/09742454A
; Patent No. US20020041876A1
; GENERAL INFORMATION:
; APPLICANT: WILEY, Steven R.
; TITLE OF INVENTION: TWEAK Receptor
; FILE REFERENCE: 2968-B
; CURRENT APPLICATION NUMBER: US/09/742,454A
; CURRENT FILING DATE: 2000-12-19
; PRIOR APPLICATION NUMBER: 60/172,878
; PRIOR FILING DATE: 1999-12-20
; PRIOR APPLICATION NUMBER: 60/203,347
; PRIOR FILING DATE: 2000-05-10
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 898
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (52)..(873)

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06:25:17 2004

us-09-245-198a-3.rnpb

ION: Description of Artificial Sequence: human TWEAK
ION: fusion protein construct

45.8%; Score 629.2; DB 9; Length 898;
arity 99.5%; Pred. No. 1.1e-162;
onservative 0; Mismatches 3; Indels 0; Gaps 0;
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GAAGAGGATCGAGCGCCATTTATGAAGTTTATCCAGCACTGGACAGGAGCG 489
CAGGTGGAGCGGACAGTGAAGTGGCTGGAGAGAGAGAGAGAGAGAGAGAG 531
CAGGTGGAGCGGACAGTGAAGTGGCTGGAGAGAGAGAGAGAGAGAGAGAG 549
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TACTCTCCAGGTTCATGAGGGGCGCTGG 865
TACTCTCCAGGTTCATGAGGGGCGCTGG 883

cation US/09883777

0110853A1

ION:

ON: Steven R.

2968-C

ION NUMBER: US/09/883,777

DATE: 2001-06-18

IN NUMBER: US 60/172,878

IN NUMBER: 1999-12-20

IN NUMBER: US 60/203,347

IN NUMBER: PCT/US00/34755

IN NUMBER: 2000-12-19

IN NUMBER: US 09/742,454

INOS: 16

In version 3.1

SEQ ID NO 1
LENGTH: 898
TYPE: DNA
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: TWEAK fusion protein construct
NAME/KEY: CDS
LOCATION: (52)..(873)
OTHER INFORMATION:
US-09-883-777-1

Query Match
Best Local Similarity 45.8%; Score 629.2; DB 9; Length 898;
Matches 631; Conservative 0; Mismatches 3; Indels 0; C

QY 232 AGTTTGGGGAGCGGGCATCGCTGTCGCGCCAGGAGCTGCCAGGAGAGAGTGGT
Db 250 AGTTTGGGGAGCGGGCATCGCTGTCGCGCCAGGAGCTGCCAGGAGAGTGGT
QY 292 GAGGAGGACGAGGACCGCTCGGAACTGAAATCCCGAGAGAGAGAGAGAGAGTCC
Db 310 GAGGAGGACGAGGACCGCTCGGAACTGAAATCCCGAGAGAGAGAGAGAGTCC
QY 352 CTTTTCCTGAACCGACTAGTTCGGCTCGCAGAGTGCATTAAGAGCGGAAAC
Db 370 CTTTTCCTGAACCGACTAGTTCGGCTCGCAGAGTGCATTAAGAGCGGAAAC
QY 412 GCTGGAAGAGGATCGCAGCGCCATTTATGAAGTTTATCCAGCACTGGACAGGAGCG
Db 430 GCTGGAAGAGGATCGCAGCGCCATTTATGAAGTTTATCCAGCACTGGACAGGAGCG
QY 472 CAGCAGGTGTGGAGCGGACAGTGAAGTGGCTGGAGAGAGAGAGAGAGTCC
Db 490 CAGCAGGTGTGGAGCGGACAGTGAAGTGGCTGGAGAGAGAGAGAGAGTCC
QY 532 CTTCTGCGCTTACAAACCGCAGATCGGGAGTGTATAGTACCCCGGCTGGCTCTA
Db 550 CTTCTGCGCTTACAAACCGCAGATCGGGAGTGTATAGTACCCCGGCTGGCTCTA
QY 592 CTGTACTGTGAGTGCATTTGATGAGGGAGAGAGTGTCTTACCTGAAGCTGGAGT
Db 610 CTGTACTGTGAGTGCATTTGATGAGGGAGAGAGTGTCTTACCTGAAGCTGGAGT
QY 652 GTGGATGTGTGTGTCGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG
Db 670 GTGGATGTGTGTGTCGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG
QY 712 CTGGGCGCGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG
Db 730 CTGGGCGCGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG
QY 772 TCCTTCGGATCGGACCGCTGCCAGTGTCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG
Db 790 TCCTTCGGATCGGACCGCTGCCAGTGTCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG
QY 832 TTCGAGTCTTCCAGGTTCACCTGAGGCGGCGCTGG 865
Db 850 TTCGAGTCTTCCAGGTTCACCTGAGGCGGCGCG 883

RESULT 11

US-09-918-995-21225

Sequence 21225, Application US/09918995

Publication No. US20030073623A1

GENERAL INFORMATION:

APPLICANT: Hyseq, Inc.

TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED

FILE REFERENCE: 20411-756

CURRENT APPLICATION NUMBER: US/09/918,995

PRIOR FILING DATE: 2001-07-30

PRIOR APPLICATION NUMBER: US/09/235,076

PRIOR FILING DATE: 1999-01-20

GenCore version 5.1.6
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.n search, using sw model
til 7, 2004, 17:41:57 ; Search time 15.6228 Seconds
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-09-245-198A-4
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SUM62
xop 10.0 , Gapext 0.5

1366 seqs, 96191526 residues

.s satisfying chosen parameters: 283366

th: 0
th: 2000000000

nimum Match 0%
ximum Match 100%
string first 45 summaries

R 78:*
pir1:*
pir2:*
pir3:*
pir4:*

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
d by analysis of the total score distribution.

SUMMARIES

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2	1323	2	S27224	N-methyl-D-asparta
0	375	2	A75264	hypothetical prote
9	378	2	H87333	HlyD family secret
9	878	2	T17245	hypothetical prote
8	206	2	T34961	probable membrane
7	441	2	S41710	mitosis-specific c
5	776	2	T36946	probable cation-tr
5	1657	2	T15838	hypothetical prote
4	310	2	D70745	hypothetical prote
4	351	2	B34768	ORF5 protein - Orf
4	566	2	T35203	probable two-compo
3	212	2	A70611	hypothetical prote
3	664	2	D83231	hypothetical prote
3	681	2	H83044	2,4-dienoyl-CoA re
3	1217	2	T00270	hypothetical prote
3	814	2	G02878	cadherin-15 precu
3	1329	2	D87226	conserved hypothet
2	660	2	T03038	probable inhibitor
2	886	2	S07132	hypothetical prote
1	210	2	D87394	hypothetical prote
1	278	2	A49266	fas ligand - rat
1	281	2	I38707	Fas ligand - human
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1	531	2	C83153	conserved hypothet
1	492	2	A87471	hypothetical prote
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0	204	1	S24641	lymphotoxin - bovi
0	445	2	T30604	hypothetical prote

30	87	6.0	2124	2	H83357	probable nc
31	86.5	6.0	399	2	F83633	hypothetical
32	86.5	6.0	439	2	E70629	hypothetical
33	86.5	6.0	755	2	E75346	probable co
34	86.5	6.0	762	2	E98121	hypothetical
35	86.5	6.0	764	1	S14113	1-phosphati
36	86.5	6.0	810	2	D95256	ATP-depende
37	86.5	6.0	887	2	AG0535	ClpB-like p
38	86	6.0	139	2	B69953	hypothetical
39	86	6.0	565	2	G98331	probable ol
40	86	6.0	568	2	E83325	probable ch
41	86	6.0	777	2	A87309	hypothetical
42	86	6.0	839	2	F75518	hypothetical
43	86	6.0	933	1	B48349	glycoprotei
44	85.5	5.9	563	2	AE3059	hypothetical
45	85.5	5.9	563	2	C98227	hypothetical

ALIGNMENTS

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4-1BB ligand - human
C:Species: Homo sapiens (man)
C:Date: 29-May-1998 #sequence_revision 29-May-1998 #text_change 21-Jul-
C:Accession: I38427
R:Alderson, M.R.; Smith, C.A.; Tough, T.W.; Davis-Smith, T.; Armitage,
Eur. J. Immunol. 24, 2219-2227, 1994
A:Title: Molecular and biological characterization of human 4-1BB and i
A:Reference number: I38426; MUID:94374434; PMID:8088337
A:Accession: I38427
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-254 <RES>
A:Cross-references: EMBL:U03398; NID:G571322; PIDN:AAA53134.1; PID:G571

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Best Local Similarity 26.5%; Pred. No. 0.32;
Matches 74; Conservative 31; Mismatches 93; Indels 81; Ga

QY	32	PPAPMAARRSRRRGEGPTALIVPLALGLALACGLGLLAVVSL-GRASL-S
Db	16	PPAP-----RARACRVLP-WALVAGLLLLLLAAACAVFLACPWAVSGARASPGS
QY	90	PAQELVAEEEDDSSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAAAHY
Db	68	PLRLRE-----GPFLSP-----DDPAGLLDL-----RQGMFAQLVAQNV
QY	150	PRPGDGAQAGVDGTVSGWEB---ARINSSPLRYNRQIGEFIVTRAGLYLYCQVH
Db	103	-----IDFPLSWSPDGLAGVSLTGGLSYKEDTKELWAKAGVYVFFQ--
QY	207	GKAVYLKLDLLVDG-----VLALRCLEFSATAASSILGPQLRLCQVSG-----
Db	147	-----LELRRVAGEGSGSVSLHLQLPLRSAGAGAAALATVDLPASSEARNSAFG
QY	250	-LLALRPGSSIRITLPAHLKAAPFL-----TYFGLFQV 283
Db	202	RLHLASQRLGVLHTEARAHAWQLTGATVLGLFRV 240

RESULT 2
S27224
N-methyl-D-aspartate receptor epsilon-4 chain - mouse
C:Species: Mus musculus (house mouse)
C:Date: 25-Feb-1994 #sequence_revision 01-Sep-1995 #text_change 17-Mar-
C:Accession: S27224
R:Ikeda, K.; Nagasawa, M.; Mori, H.; Araki, K.; Sakimura, K.; Watanabe,
FEBS Lett. 313, 34-39, 1992
A:Title: Cloning and expression of the epsilon-4 subunit of the NMDA rec
A:Reference number: S27224; MUID:93050214; PMID:1385220
A:Accession: S27224

us-09-245-198a-4.rpr

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QY 25 GAVRQAQPPAPMAARRSRR-----RGRGPGTALLVPLALGLGLALA
Db 19 GAIKOKS-----MAVEKNRRALGIDGNVTVRGEVKALPQVSRPITRGF-----
QY 74 -----LAVVLSGRASLS-----AQEPAQBELVABEDODPSLNPO
Db 70 ANAEAAAAENNNKNSLVANAKGADGALPIKRVAVRPVQKTKVSKPQETIIIESPT
QY 115 DPAPFLNLRPRKS-----APGKRKTRARRATAAHYEVHPRPQ-----DGAQAQAG
Db 130 ---APVLEKEITGKSLKKKAPTTLTSTLTARSKAASV-VTKPKQEIVIDDAVDNN
QY 166 SWEF-----ARINSSPLRY-----NROIGPIVTRAGLYLYCQVHFD---EG
Db 187 VEYVEDMYKFKYSAENDSRPHDYMDSQPEINEKM--RAILIDLWLVQVHVKPELSE
QY 213 KLDLLVDGVIALRC-----LFEFSATASSIGPOLRLCQVSGLLALRPGS 257
Db 245 TIN-IVRYLASKTTSRRELQILGMSMLIASKYBEIMAPEVNDLVCSIDGS 295

RESULT 8
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C:Species: Streptomyces coelicolor
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 18-Aug
C:Accession: T36946
R:Seeger, K.J.; Harris, D.; Thomson, N.R.; Parkhill, J.; Barrell, B.G.
submitted to the EMBL Data Library, September 1999
A:Reference number: 221607
A:Accession: T36946
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 1-776 <SEE>
A:Cross-references: EMBL:AL109862; PIDN: CAB53131.1; GSFPD: GN00070; SCO
A:Experimental source: strain A3(2)
C:Genetics:
A:Gene: SCORDB: SCJ1.13
C:Superfamily: ATPase nucleotide-binding domain homology
F:442-585/Domain: ATPase nucleotide-binding domain homology <ATN>

Query Match 6.5%; Score 93.5; DB 2; Length 776;
Best Local Similarity 24.1%; Prd; No. 12;
Matches 63; Conservative 30; Mismatches 103; Indels 65; G; G;

QY 47 RRGEPTGTVLPLALGLGLALA--CLGLLLA-VVSLGSRASLSAQEPAQBELVAEE
Db 73 RRHGAGVDLIVLALUGGTLVAGEYIAGVLIALMLATGRTLEGAAQRASHDLHALL
QY 104 SELNPQTESQDPAPPLNLRPRRSAPKGRKTRARRATAAHYEVHPRPQDGAQA
Db 133 RSARRRTGDG-----VVR-----VPLSEITAGDALVVGPEVVP-----
QY 164 TVSGWE---EARINSSPLRYNRQIGE----FIVTRAGLYLYCQVHFDGKAVYLI
Db 170 RVESTEAVILDESVLCEPIQVTRQREGARGSAVAGGAFDL-----RATAII
QY 217 LVQGVIALRCLEFSATASSIGPOLRLCQ-----VSGLLALRPGSSLR
Db 221 TYAGIVRL-----AQQAGASAPVVRADRYAAWFLPLATAALAWLVSGSAVFA
QY 261 -IRTLFWHLKKAAPFLTYFGL 280
Db 275 LVVATPCPILLAAPVAVVSGL 295

RESULT 9
T115838
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C:Species: Caenorhabditis elegans
C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 17-Mar-
C:Accession: T115838
C:Minx, P.

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EMBL Data Library, October 1995
 e sequence of *C. elegans* cosmid C54D2.
 r: Z18415
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 nary; translated from GB/EMBL/DBJ
 DNA
 7 <MIN>
 9: EMBL:U37548; NID:G1017804; PID:G1017809; PIDN:AAA79201.1; CESP:C54D2
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 dium channel protein
 6.5%; Score 93.5; DB 2; Length 1657;
 larity 23.2%; Pred. No. 29;
 Conservative 38; Mismatches 99; Indels 71; Gaps 12;
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 QTNPAALYFVALMTFGNYVLFNLLVAILVEGQESKEEKRLQLEEDARKQAVEED 940
 SELNPQTESQDPAPFLNLRVP-----RRSAPKGRKTRARRAIAAHYEVHPRPGQ 154
 RELELLIAKTSPA--FNNGVAPAECTCQRPSP--ESSPRLLSANY--HSPER 994
 QAGVDGTVSGWEARINSSPL-----RYNRQIGEFIVTRAGLYLYCQV 202
 -ANLDAIID--KRLVLRNSAPFDRSPVSEGRDSDRLNRHASLVLVPVANGVYRQRV 1051
 -----FDEGKAVVLLDLVDGVALRCLE---ESATAASLGLFQQLRLCQV 247
 KASQELKQALAEKKEAKNEAKQNTFVRKLLKTKLHNTEFS----- 1095
 LALRPGSSIRITLPWAHLKAAPFLTYF 278
 FLMPKPNLRKIKLQTTQKKWEDYVLF 1124
 in Rv0497 - Mycobacterium tuberculosis (strain H37RV)
 erium tuberculosis
 98 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
 15
 sch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.
 tes, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.
 Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 14, 1998
 R.; Suleston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 ng the biology of Mycobacterium tuberculosis from the complete genome
 : A70500; MUID:98295987; PMID:9634230
 15
 lary; nucleic acid sequence not shown; translation not shown
 INA
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 : GB:Z77162; GB:AL123456; NID:G3261606; PIDN:CAB00923.1; PID:e255036;
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 arity 24.4%; Pred. No. 4.8;
 onservative 25; Mismatches 103; Indels 120; Gaps 18;
 :LSGSRDGA---VRQAQPPAPMAARRSQRRG-----R 47
 :SSGNQISVAELLARQGTGAP--ARRRRRRGSDAITVAELTGETPIIRDHHH 63
 GTALLVPLALGLGALACLGLLAVVSLGSRASLSAQEPAQEELVAEE----- 99
 AHASQSPAANGR-----VQGEAAPQSPAEPPVAEQ-VAEPEFTTVVWS 109
 SELNPQTESQDPAPFLNLRVP-----SAPKGRKTRARRAI---AAHY----- 146

Db 110 QPERWPKSPQDRRESGPELSEYPRRLRTHSDRAPAGPPSGAEHMSPPDVEH:
 QY 147 -----EVHPRPGQDG-----AQAGVDGTVSGWEARINSSS-----
 Db 170 DVLDTVEGEAEATEVREAQGRGERHAAAGAAAGTVEGDGAARVARALDVI
 QY 183 RQIGEEIVTR-----AGLYLYCQVHFE-----GKAVYLLKDL-----LVDC
 Db 230 ---GALVLIOSILAVAFGAGLF-----IAFDQLWRMNSIVALVLSVMVILGLVVS
 QY 226 CLBEFSAT-----AASSLGPQLRLCQ 246
 Db 282 KTDIASTLIAVAVGALITLGP-LALLQ 308
 RESULT 11
 B34768
 ORF5 protein - Orf virus (strain NZ2)
 C;Species: Orf virus
 C;Date: 23-Aug-1991 #sequence_revision 23-Aug-1991 #text_change 08-Oct-1991
 C;Accession: B34768
 R;Fraser, K.M.; Hill, D.F.; Mercer, A.A.; Robinson, A.J.
 Virology 176, 379-389, 1990
 A;Title: Sequence analysis of the inverted terminal repetition in the
 A;Reference number: A34768; MUID:90266454; PMID:2129563
 A;Accession: B34768
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-351 <FRA>
 A;Cross-references: GB:M30023; EMBL:M37623; NID:G332561; PIDN:AAA4678
 Query Match 6.4%; Score 92; DB 2; Length 351;
 Best Local Similarity 23.0%; Pred. No. 6.6;
 Matches 45; Conservative 18; Mismatches 73; Indels 60;
 QY 14 PLPRSLGSRDGGAVQAQAPPAARRSQ-RRGRGEGPTALLVPLALGLGAL
 Db 211 PLPRRAAR---GQRGQPPPRARRAQPRRRAPRAAG-----
 QY 73 LLAVVSLGSRASLSAQEPAQEELVAEEDQPSLENPQTESQDPAPFLNLRVPR
 Db 248 -----ARRGRGAPRQOQRPVQRAAAAQRRRAQOR
 QY 133 GKTRARRAIAHYE-VHPRQDGAQAGVDGTVSGWEARINSSPLRYNRQIG:
 Db 284 PRVVEARRARRQORAHQR--RRGRARRTRCSTS-----RVVSKD-----SREVG
 QY 192 RAGLYLYCQVHFEFG 207
 Db 333 KERYIRRVLLHFEFG 348
 RESULT 12
 T35203
 probable two-component sensor - Streptomyces coelicolor
 C;Species: Streptomyces coelicolor
 C;Date: 05-Nov-1999 #sequence_revision 05-Nov-1999 #text_change 31-Jan-1999
 C;Accession: T35203
 R;Seeger, K.; Harris, D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.J.
 submitted to the EMBL Data Library, April 1998
 A;Reference number: Z21571
 A;Accession: T35203
 A;Status: preliminary; translated from GB/EMBL/DBJ
 A;Molecule type: DNA
 A;Residues: 1-566 <SEE>
 A;Cross-references: EMBL:AL02374; PIDN:CAA18527.1; GSPDB:GN00070; SCC
 A;Experimental source: strain A3(2)
 C;Genetics:
 A;Gene: SCOEDB:SC5B8.19C
 C;Superfamily: two-component sensor histidine kinase; sensor histidine
 Query Match 6.4%; Score 92; DB 2; Length 566;

arity 24.5%; Pred. No. 11;
 conservative 24; Mismatches 99; Indels 80; Gaps 14;
 MAARRSQR-----RGRGEP-----GTALLVPLALGLALACGLLLAVVSLGS-- 81
 RTVMAGSTTPVRLRLGLPRVFSQVLLMQLAAGVAVLATGLFLA--PLGDQL 59
 --RASLSAQEPAQBELVAEBDDQPSLNPQTESQDPAPFLNLRVPRRSAPKGRK 135
 MRRLAIAQTAAQPPQVVRD-----LRTTRTPANGPVQRE 98
 PRATAAHYEV-----HPRPCDQAQAGVD--GTVSGWEEARINSSS----- 177
 REATRAEYVVMMDRQGVNWSHTDPERIGEVVSTDPQALACREVMEDDGLGRSA 158
 PLRYNRQIGEFI-VTRAGLYLYLCO---VHFDEGKAVY-----LKLDDLVDGVLAR 225
 PLRDGD--GEIVGAVSVGIADSVARLIHAIPGLFAYAGGALAVGALASWIIISR 216
 IFSATAASSLGPQLRLCQVSGLLALR 254
 YTFDLAFS-----DIAGLLAER 236

ain Rv1219c - Mycobacterium tuberculosis (strain H37RV)
 atrium tuberculosis
 98 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
 11
 sch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.
 ies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.
 Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 44, 1998
 ing R.; Suleston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 ing the biology of Mycobacterium tuberculosis from the complete genome
 r: A70500; MUID:98295987; PMID:9634230
 11
 nary; nucleic acid sequence not shown; translation not shown
 DNA
 <COL>
 s: GB:Z93777; GB:AL123456; NID:G3261726; PIDN:CAB07841.1; PID:ei299832;
 urce: strain H37RV

6.3%; Score 91.5; DB 2; Length 212;
 larity 24.5%; Pred. No. 4.1;
 Conservative 26; Mismatches 78; Indels 59; Gaps 9;
 RGPPTALLVPLALGLL-----ALACGLLLAVVSLGSASLSAQEPAQBELVAEB-- 99
 IG-----FGVGLRAIAEAGVSAALVIHFHFSKEGL---RKACDDFVAEEIR 66
 -----DQDPSLNPQTESQDPAPFLNLRVPRRSAPKGRKTRARRAJAAHYEVHPR 151
 KAAALKENDPTTTLAQMAETSAEAPLMAYLVRSMSQSGELAKMLWQKI----- 117
 QDGAQAGVDGTVSGWEEARINSSSPLRYNRQIGEFI-VTRAGLYLYLCOVHFD----- 205
 -DNAEEYLD-----EGVRAGTVKPSRDPARARFLAITGGGFFLLYLMHENPTDLR 168
 -----EGKAVYLKDLLVDGVLALCL--EFSATA 234
 LRDYAHDMVLPSEVYVTEGLLADRAVMEAFLLAE 204

tein PA3305 [imported] - Pseudomonas aeruginosa (strain PA01)
 monas aeruginosa
 :000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
 .231

R;Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.
 adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.
 .; Lory, S.; Olson, M.V.
 Nature 406, 959-964, 2000
 A>Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an c
 A;Reference number: A82950; MUID:20437337; PMID:10984043
 A;Accession: D83231
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-664 <STO>
 A;Cross-references: GB:AE004753; GB:AE004091; NID:G9949433; PIDN:AAG06
 A;Experimental source: strain PA01
 C;Genetics:
 A;Gene: PA3305

Query Match 6.3%; Score 91.5; DB 2; Length 664;
 Best Local Similarity 25.8%; Pred. No. 15;
 Matches 77; Conservative 28; Mismatches 117; Indels 77;
 QY 28 RQAQPPAPMAARR-----SOR-----RGRGEPGTALLVPLALGLALACI
 Db 194 RQRPQGLLNALSKIVEVDAQRDHAHFEGERRRAGALALLSRDLISL-
 QY 75 AVVSLGSASLSAQEPAQEE-----LVAEEDQDPSLNPQTESQDPAPFLNRL
 Db 246 ARGVARQARLSEEEERRVERVWLAALASALEGTDPASMQALRELAQVA-----
 QY 129 SAPKGRKTRARRAJAAHYEVHPRPQDGAQAGVDGTVSGWEEARINSSSPLRYNR
 Db 300 SNDQ-RYLLTRCSVLLKAVN-----AEKGMRAVASGEVGRVGSAGTLSWHR
 QY 187 EFIVTPAGL-----YLYL-----COVHFDGKAVYLKDLLV
 Db 352 LFYGTSSALALLGLSVYVIYTAWPAASGAMLLAAVVCSEFANPDNAVIGLSPLR
 QY 224 LRCLFEFSATAASSLGPQLRLCQVSG--LIALRPGSSLRIRTPMAHLKAPFLI
 Db 412 I-----PAAMLVISQWLLPQWNGFPLICLAWGVLPFPATLGMVFPVTAGTAI

RESULT 15
 H83044
 2,4-dienoyl-CoA reductase PadH2 PA4814 [imported] - Pseudomonas aeru
 C;Species: Pseudomonas aeruginosa
 C;Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-D
 C;Accession: H83044
 R;Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.
 adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, J
 .; Lory, S.; Olson, M.V.
 Nature 406, 959-964, 2000
 A>Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an
 A;Reference number: A82950; MUID:20437337; PMID:10984043
 A;Accession: H83044
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-661 <STO>
 A;Cross-references: GB:AE004894; GB:AE004091; NID:G9951076; PIDN:AAG
 A;Experimental source: strain PA01
 C;Genetics:
 A;Gene: fadh2; PA4814
 C;Superfamily: Methylophilus methylotrophus W3A1 trimethylamine dehy

Query Match 6.3%; Score 91; DB 2; Length 681;
 Best Local Similarity 26.3%; Pred. No. 17;
 Matches 79; Conservative 31; Mismatches 86; Indels 104;
 QY 6 FEISARRPLPRSLGRDGGAVRQAQPPAP-----MAARRSQRRGR-----
 Db 442 FRVRLERLGVLDLGRH-----VRQELDQDFDDVVVATGIPRRPRIDGIGPT
 QY 48 --RGEPTALLVPLALGLALACGLLLAVVSLGSASLSAQEPAQBELVAE
 Db 498 VLRGAPVGARVAIVGAGGIGFDVA--AFLVAAPSDG-----QPRALGEWLAI

6:25:24 2004

us-09-245-198a-4.rspt

GenCore version 5.1.6
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1 search, using sw model

il 7, 2004, 17:41:27 ; Search time 39.6149 Seconds
(without alignments)
2261.954 Million cell updates/sec

09-245-198A-4
4
SLLDPEISARRLPRLSLG.....PWAHLKAAPFLTYFGLFQVH 284

SUM62

op 10.0 , Gapext 0.5

7041 seqs, 315518202 residues

s satisfying chosen parameters: 1017041

th: 0

th: 2000000000

nimum Match 0%

ximum Match 100%

string first 45 summaries

TREMBL_25:*

sp archaea:*

sp bacteria:*

sp fungi:*

sp human:*

sp invertebrate:*

sp mammal:*

sp mhc:*

sp organelle:*

sp phage:*

sp plant:*

sp rodent:*

sp virus:*

sp vertebrate:*

sp unclassified:*

sp virus:*

sp bacteriaph:*

sp archaea:*

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
d by analysis of the total score distribution.

SUMMARIES

xy	ch	Length	DB	ID	Description
9	410	11	Q8BXS2	Q8bxs2 mus musculus	
10	330	4	Q81zk7	Q81zk7 homo sapien	
11	261	5	Q8MRW2	Q8mrw2 drosophila	
12	325	5	Q9V5G2	Q9v5g2 drosophila	
13	409	5	Q8MY88	Q8my88 drosophila	
14	415	5	Q8MUJ1	Q8muj1 drosophila	
15	409	5	Q8IGD3	Q8igd3 drosophila	
16	398	6	Q8MK49	Q8mk49 sorex ciner	
17	532	16	Q82HP6	Q82hp6 streptomyc	
18	565	16	Q8KY66	Q8ky66 streptomyc	
19	955	10	Q84T85	Q84t85 oryza sativ	
20	967	10	Q7X1L4	Q7x1l4 oryza sativ	
21	643	16	Q9KZ17	Q9kz17 streptomyc	
22	375	16	Q9RRH5	Q9rrh5 deinococcus	
23	893	16	Q81ZX0	Q81zx0 streptomyc	
24	611	11	Q8KOM8	Q8kom8 mus musculus	

17	100	6.9	850	11	Q9JJ15	Q9jj15 mus
18	100	6.9	850	11	Q8OXI6	Q8oxi6 mus
19	99.5	6.9	378	16	Q9AAB9	Q9aab9 caul
20	99	6.9	330	4	Q8N5L1	Q8n5l1 homo
21	99	6.9	614	4	Q7Z4K2	Q7z4k2 homo
22	99	6.9	915	4	Q7Z5I1	Q7z5i1 homo
23	98	6.8	206	16	Q9S2W5	Q9s2w5 stre
24	97.5	6.8	694	16	Q82FL1	Q82fl1 stre
25	97.5	6.8	1560	4	Q96JP2	Q96jp2 homo
26	97	6.7	408	10	Q8S5I5	Q8s5i5 oryz
27	97	6.7	926	4	Q9NYA0	Q9nya0 homo
28	97	6.7	1058	4	Q9Y4G2	Q9y4g2 homo
29	96.5	6.7	975	11	Q8BWB1	Q8bwb1 mus
30	96	6.6	629	10	Q8SIA6	Q8sia6 oryz
31	95.5	6.6	536	4	Q9HB96	Q9hb96 homo
32	95.5	6.6	655	16	Q9FBR7	Q9fbr7 stre
33	95	6.6	340	16	Q7WFI3	Q7wfi3 bord
34	95	6.6	390	2	Q8KW28	Q8kw28 ruege
35	95	6.6	810	16	Q82K60	Q82k60 stre
36	94.5	6.5	748	5	Q8T2Y0	Q8t2y0 trypa
37	94.5	6.5	1696	11	Q9WTR8	Q9wtr8 ratt
38	94	6.5	340	16	Q7W3N7	Q7w3n7 bord
39	94	6.5	340	16	Q7W0H0	Q7w0h0 bord
40	94	6.5	448	16	Q886A1	Q886a1 pseu
41	94	6.5	937	16	Q93JD1	Q93jd1 stre
42	94	6.5	1910	10	Q7XU19	Q7xu19 oryz
43	93.5	6.5	776	16	Q9RJ01	Q9rj01 stre
44	93.5	6.5	854	16	Q9F2P0	Q9f2p0 stre
45	93.5	6.5	1038	10	Q9AS09	Q9as09 oryz

ALIGNMENTS

RESULT 1

Q8BXS2	ID	Q8BXS2	PRELIMINARY;	PRT;	410 AA.
AC	Q8BXS2;				
DT	01-MAR-2003	(TREMBLrel. 23, Created)			
DT	01-MAR-2003	(TREMBLrel. 23, Last sequence update)			
DT	01-OCT-2003	(TREMBLrel. 25, Last annotation update)			
DE	Tumor necrosis factor.				
OS	Mus musculus (Mouse).				
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
OX	NCBI_TaxID=10090;				
RN	[1]_TaxID=10090;				
RP	SEQUENCE FROM N.A.				
RC	STRAIN=C57BL/6J; TISSUE=Retina;				
RX	MEDLINE=22354683; PubMed=12466851;				
RA	The FANTOM Consortium,				
RA	the RIKEN Genome Exploration Research Group Phase I & II Team;				
RT	"Analysis of the mouse transcriptome based on functional annotation"				
RT	60,770 full-length cDNAs."				
RL	Nature 420:563-573 (2002).				
DR	EMBL; AK044387; BAC31897.1;				
DR	PIR; PT0714; PT0714.				
DR	GO; GO:0016020; C:membrane; IEA.				
DR	GO; GO:0005164; F:tumor necrosis factor receptor binding; IEA.				
DR	GO; GO:0005955; P:immune response; IEA.				
DR	InterPro; IPR006052; TNF_family.				
DR	InterPro; IPR008983; TNF_like.				
DR	SMART; SM00207; TNF; 2.				
DR	PROSITE; PS00251; TNF 1; 1.				
DR	PROSITE; PS50049; TNF 2; 2.				
SQ	SEQUENCE 410 AA; 45881 MW; 590A4B74C33FB8D4 CRC64;				

Query Match 64.9%; Score 937.5; DB 11; Length 410;
Best Local Similarity 79.1%; Pred. No. 1.3e-73;
Matches 193; Conservative 11; Mismatches 29; Indels 11; G

QY

36 MAARRQRGRGRGPGTALLVPLALGLALACLLAVVSLGSRASLSA-QEP;
|||||

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us-09-245-198a-4.rsp

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RSRRRRRRGEGTALLAPLVLSGLALACLLGLLVVSLGSWATLSAQEPFSOBE 60
EDDDPSLNPTQTESQDPAPFLNRLVRRPSAPKGRKTRARRAIAAHYEVHPRPQ 154
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
3DRPEPELNPTQTESQDVVPFLEQLVRRPSAPKGRKTRARRAIAAHYEVHPRPQ 120
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
JAGVDGTVSGWEAPARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDGKAVYLK 214
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
JAGVDGTVSGWEETKINSSPLRYNQIGEFIVTRAGLYLYCQVHFDGKAVYLK 180
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
JGVLAIRCLESFASATAASGLGQLRLCQVSGLLALR-----PGSLRLRITLP 265
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
JGVLAIRCLESFASATAASPGQLRLCQTE-LQSLRRVRSRLQRSGPQKQGERP 239
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269
243
PRELIMINARY; PRT; 330 AA.
(TMREBLrel. 23, Created)
(TMREBLrel. 23, Last sequence update)
(TMREBLrel. 25, Last annotation update)
(Human)
stazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Primates; Catarrhini; Hominidae; Homo.
506;
4 N.A.
9924; PubMed=12411489;
B.; Medema J.P., Lopez-Fraga M., Lozano J.C.,
M., Picard A., Martinez-A C., Garcia-Sanz J.A.,
is hybrid mRNA encodes TWE-PRIL, a functional cell surface
fusion protein."
11-5720(2002).
1; AAL90443.1;
20; C:membrane; IEA.
34; F: tumor necrosis factor receptor binding; IEA.
35; P: immune response; IEA.
3006052; TNF family.
3008983; TNF_like.
3; TNF; 1.
37; TNF; 1.
3251; TNF; 1.
3049; TNF; 2.
30 AA; 36588 MW; FC6F3BCA29C029AE CRC64;
Larity 58.4%; Score 844; DB 4; Length 330;
Conservative 100.0%; Pred. No. 1.5e-65; Indels 0; Gaps 0;
RSRRRRRRGEGTALLAPLVLSGLALACLLGLLVVSLGSRASLSAQEPQEL 95
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
RSRRRRRRGEGTALLAPLVLSGLALACLLGLLVVSLGSRASLSAQEPQEL 60
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
EDDDPSLNPTQTESQDPAPFLNRLVRRPSAPKGRKTRARRAIAAHYEVHPRPQ 155
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
EDDDPSLNPTQTESQDPAPFLNRLVRRPSAPKGRKTRARRAIAAHYEVHPRPQ 120
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
AGVDGTVSGWEAPARINSSPLRYNQIGEFIVTRAGLYLYCQ 201
: : : : : : : : : : : : : : : : : : : : : : : : : : : :
AGVDGTVSGWEAPARINSSPLRYNQIGEFIVTRAGLYLYCQ 166
PRELIMINARY; PRT; 261 AA.

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DT 01-OCT-2002 (TMREBLrel. 22, Created)
DT 01-OCT-2002 (TMREBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TMREBLrel. 25, Last annotation update)
DE SD18286P.
GN EIGER OR CG12919.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
RN [1]
SEQUENCE FROM N.A.
RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
RA Champe M., Chavez C., Dorsett V., Dresnek D., Farfan D., Frise E
RA George R., Gonzalez M., Guarin H., Kronmiller B., Li P., Liao G.,
RA Miranda A., Mungall C.J., Nunoo J., Pacleb J., Paragas V., Park
RA Patel S., Phouanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M
RA Celniker S.;
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY119233; AAM51093.1; -.
DR FlyBase; FBgn0033483; eiger.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0005164; F:tumor necrosis factor receptor binding; IEA.
DR GO; GO:0006955; P:immune response; IEA.
DR InterPro; IPR006052; TNF family.
DR InterPro; IPR008983; TNF_like.
DR SMART; SM00207; TNF; 1.
DR PROSITE; PS00251; TNF_1; 1.
DR PROSITE; PS50049; TNF_2; 1.
DR PROSITE; PS50049; TNF_2; 1.
SQ SEQUENCE 261 AA; 29780 MW; 13B6D5A04EC9122C CRC64;
Query Match 8.0%; Score 116; DB 5; Length 261;
Best Local Similarity 25.4%; Pred. No. 0.048;
Matches 57; Conservative 32; Mismatches 97; Indels 38;
QY 82 RASLSAQEPQAEELVAE-----EDQPSLNPTQTESQDPAPFLNR-----
DB 54 RKSRSIADVRNEEQNIQGNHTELOEKSNEATSKES--PAPLHRRRMRHRL
QY 128 RSAPKGRKTRARRAIAAHYEVHPRPQDCAQAGVDGTVSGWEAPARINSSPLRYN
DB 112 ESSLARSSEDSRP--AAHFLSSRRRHQSGM-GYHGDVYIGNDNERNYSQG-HFQ
QY 188 FIVTRAGLYLYCQV-----HFDEGKAVYLKDLLVDGVLARCLEEFSAATASS
DB 168 LTVTNTGLVYVYAQICYNNSHDQGFIVE-----QGDTPLQCLN---TVPTN
QY 243 RLCQVSGLLALRPGSSLRITL---PWAHLKAAPFLTYGFLFQV 283
DB 218 HTCHTSGLIHLERNRIHLKDIHNDRNAVLREGNRSYFGIFKV 261
RESULT 4
Q9V5G2
ID Q9V5G2 PRELIMINARY; PRT; 325 AA.
AC Q9V5G2;
DT 01-MAY-2000 (TMREBLrel. 13, Created)
DT 01-MAY-2000 (TMREBLrel. 13, Last sequence update)
DT 01-OCT-2003 (TMREBLrel. 25, Last annotation update)
DE CG12919 protein.
GN EIGER OR CG12919.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
RN [1]
SEQUENCE FROM N.A.
RC STRAIN=Berkeley;
RX MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.)

```

01-OCT-2002 (TReMBLrel. 22, Last sequence update).	
01-OCT-2003 (TReMBLrel. 25, Last annotation update)	
TNF superfamily ligand, Eiger (Tumor necrosis factor family member Dtl).	
EIGER OR Dtl OR CG12919.	
Drosophila melanogaster (Fruit fly).	
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.	
NCBI_TaxID=7227;	
[1]	
SEQUENCE FROM N.A.	
MEDLINE=22060500; PubMed=12065414;	
Igaki T., Kanda H., Yamamoto-Goto Y., Kanuka H., Kuranaga E., Aigaki T., Miura M.;	
"Eiger", a TNF superfamily ligand that triggers the Drosophila JNK pathway.";	
EMBO J. 21:3009-3018 (2002).	
[2]	
SEQUENCE FROM N.A.	
Inohara N., Nunez G.;	
"Dtl", a Drosophila tumor necrosis factor family member.";	
Submitted (MAY-1999) to the EMBL/GenBank/DBJ databases.	
EMBL; AB073865; BAC00950.1;	
EMBL; AF149799; AAC01510.1;	
FlyBase; FBgn0033483; eiger.	
GO; GO:0016020; C:membrane; IEA.	
GO; GO:0005164; F:tumor necrosis factor receptor binding; IEA.	
GO; GO:0006955; P:immune response; IEA.	
InterPro; IPR06052; TNF family.	
InterPro; IPR008983; TNF_like.	
SMART; SM00207; TNF; 1.	
PROSITE; PS00251; TNF 1; 1.	
PROSITE; PS00049; TNF 2; 1.	
SEQUENCE 409 AA; 46331 MW; 8306AECCEL4397B8 CRC64;	
Query Match 8.0%; Score 116; DB 5; Length 409;	
Best Local Similarity 25.5%; Pred. NO. 0.084;	
Matches 56; Conservative 33; Mismatches 95; Indels 36; G	
QY	82 RASLSAQEPAQELVAE---EDQDPSELNPQTESQDPAPFLNLRVPRSPAKGI
DBD	208 RKSRSIADVNSEQNIQGMHTLQEKSNKESNEATSKES--PAPLHHR-----RMMSRHH
QY	138 AR3A-----IAAHVEVHPRPQDGAQAGVDGTGVSQWGEARINSSSPLYRNQOIGE.
DBD	262 VRKARSDSRPAAHFHLSSRRRHQSGM-GYHGDVMYIGNDNERNYSQG-HFQTRDGV.
QY	192 RAGLYLYLQYV-----HFDEGKAVYKLKDLLVDGVLAIRCLCEFSATASSLGPQL
DBD	320 NTGLIYVYVYQICVNNSHDQNGFIVF-----QGDTPFLQCLN----TVPNMPHKVI
QY	247 VSGLLALPCSSSLRIURLT---PWAHLKAAPFLTVFGLFOV 283
DBD	370 TSGLIHLERNRIHLKDHNDRNAVREGNRRSIFGIFKV 409
RESULT 6	
Q8MUJ1	PRELIMINARY; PRT; 415 AA.
IC Q8MUJ1	
AD Q8MUJ1	
DT 01-OCT-2002 (TReMBLrel. 22, Created)	
DT 01-OCT-2002 (TReMBLrel. 22, Last sequence update)	
DT 01-OCT-2003 (TReMBLrel. 25, Last annotation update)	
DE Eiger (DARTH).	
DE Eiger OR CG12919 OR DARTH.	
OS Drosophila melanogaster (Fruit fly).	
OS Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.	
OC NCBI_TaxID=7227;	
OX	[1]
RP	SEQUENCE FROM N.A

923; PubMed=12176339;
n M., Basler K.;
TNF Signaling Mechanisms. JNK-Dependent Apoptosis
Eiger, the Drosophila Homolog of the TNF Superfamily.";
2:1263-1268(2002).
N.A.
938; PubMed=12894227;
Maaty W.S., Chen P., Tomar R.S., Eby M.T., Chapo J.,
ore N., Zachariah S., Sinha S.K., Abrams J.M.,
s receptor, Wengen, comprise a TNF-like system in
860-4867(2003).
6; AAM76710.1; -
1; AAM66763.1; -
0033483; eiger.
0; C:membrane; IEA.
4; P:tumor necrosis factor receptor binding; IEA.
5; P:immune response; IEA.
006052; TNF family.
008983; TNF-like.
7; TNF; 1.
251; TNF 1; 1.
049; TNF 2; 1.
5 AA; 46918 MW; E087A26DE222D2BF CRC64;
8.0%; Score 116; DB 5; Length 415;
arity 25.4%; Pred.No. 0.086; 97; Indels 38; Gaps 10;
onservative 32; Mismatches

SAQPAQELVAE---EDQPSSELPOTERSQDPAPFLNR-----LVRPR 127
SIADVRNEEQNIQNHTELOEKSNEATSKES--PAPLHRRMHSRHLVRKG 265
KGRTRARRATAAHYEVHPRPGQGAQAGVDGTSGWEARINSSPLRYNRQIGE 187
SARSEDSP--AAHFLSSRRHQSGM-GYHGDYIGNDNRNSYQG-HFQTRDGV 321
RAGYLYLCQV-----HFDEKAVYKLDLLVDGVLALRCLEFSATASLSGQL 242
NTGLYYVYAQICYNNSHDQGFIVF-----QGDTPFLQCLN----TPTNMPHKV 371
VSGLLALRPGSSSLRIRTL---PWAHLKAAPFLTYFGLFQV 283
TSGLIHLENERIHLKDIHNDNAVLREGNRSYFGIFKV 415
RELIMINARY; PRT; 409 AA.
TREMBlrel. 23, Created)
TREMBlrel. 23, Last sequence update)
TREMBlrel. 25, Last annotation update)
lanogaster (Fruit fly).
tazaa; Arthropoda; Hexapoda; Insecta; Pterygota;
lopterygota; Diptera; Brachycera; Muscomorpha;
Drosophilidae; Drosophila.
1 N.A.
Brokstein P., Hong L., Agbayani A., Carlson J.,
iavez C., Dorsett V., Dresnek D., Farfan D., Frise E.,
nzalez M., Guarin H., Kronmiller B., Li P., Liao G.,
ungall C.J., Nunoo J., Pacleb J., Paragas V., Park S.,
uanenavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,
V-2002) to the EMBL/GenBank/DBJ databases.
18; AAN71595.1; -

DR FlyBase; FBgn0064801; BcDNA:RH51659.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0005164; F:tumor necrosis factor receptor binding; IEA.
DR GO; GO:0006955; P:immune response; IEA.
DR InterPro; IPR006052; TNF family.
DR InterPro; IPR008983; TNF-like.
DR SMART; SM00207; TNF; 1.
DR PROSITE; PS00251; TNF 1; 1.
DR PROSITE; PS50049; TNF 2; 1.
SQ SEQUENCE 409 AA; 46401 MW; FC2E9BD9E012D257 CRC64;
Query Match 7.8%; Score 113; DB 5; Length 409;
Best Local Similarity 23.7%; Pred.No. 0.15;
Matches 54; Conservative 36; Mismatches 94; Indels 44; G

QY 84 SLQAQPAQEL-----VAEEDQPSSELPQTEE-----SQDPAPFLNLVR
Db 198 SYNAAKKKKKRSKRSIADVRNEEQNIQNHTELOEKSNEATSKERAPLHRR---
QY 130 APKGRKTRARRA-----IAAHYEVHPRPGQGAQAGVDGTSGWEARINSSPL
Db 254 HSRHLLVRKARSEDSPAAHFLSSRRHQSGM-GYHGDYIGNDNRNSYQG-
QY 184 QIGEFIVTRAGLYLYLCQV-----HFDEKAVYKLDLLVDGVLALRCLEFSATP
Db 312 RDGVLTVNTGLYYVYAQICYNNSHDQGFIVF-----QGDTPFLQCLN----TV
QY 239 GPQLRLQVSGLLALRPGSSSLRIRTL---PWAHLKAAPFLTYFGLFQV 283
Db 362 PKVHTCHTSGLIHLERNERIHLKDIHNDNAVLREGNRSYFGIFKV 409

RESULT 8
Q8MK49 PRELIMINARY; PRT; 398 AA.
ID Q8MK49
AC Q8MK49;
DT 01-OCT-2002 (TrEMBlrel. 22, Created)
DT 01-OCT-2002 (TrEMBlrel. 22, Last sequence update)
DT 01-JUN-2003 (TrEMBlrel. 24, Last annotation update)
DE Alpha 2B adrenergic receptor (Fragment).
GN ADRA2B.
OS Sorex cinereus (Masked shrew).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Insectivora; Soricidae; Soricinae; Sorex.
OX NCBI_taxID=36803;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21608557; PubMed=11743200;
RA Murphy W.J., Eizirik E., O'Brien S.J., Madsen O., Scally M.,
RA Douady C.J., Teeling E., Ryder O.A., Stanhope M.J., de Jong W.W.,
RA Springer M.S.;
RA "Resolution of the early placental mammal radiation using Bayesian
RT phylogenetics";
RL Science 294:2348-2351(2001).
DR EMBL; AJ315936; CAC87000.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin.
DR InterPro; IPR000276; GPCR_Rhodopsn.
DR Pfam; PF00001; 7tm_1; 1.
DR PRINTS; PR00237; GPCR_RHODOPSIN.
DR PROSITE; PS00237; G-PROTEIN RECEPTOR FL 1; 1.
DR PROSITE; PS50262; G-PROTEIN RECEPTOR FL 2; 1.
KW Receptor.
FT NON TER 1 1
FT NON TER 398 398
SQ SEQUENCE 398 AA; 43576 MW; D57E67B689535E27 CRC64;
Query Match 7.2%; Score 104; DB 6; Length 398;
Best Local Similarity 25.2%; Pred.No. 0.91;
Matches 55; Conservative 23; Mismatches 76; Indels 64; C

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SRDGAIVRQA-QPP-APMAAR-----RSRRRRGRGEP 51
|||:||||:|
|SAGKGSQAGQAGPAGPVSAKPLSLASPLVVGADAPGLSKPPRENMETGAPEP 253
|
|PLALGLGLAL-----ACLGLLAVSLGSRASLSAQEPAQBELVAEEDDPSE 105
|PSPWPAFPSAODLKEGACV-----VSIEEFAEEDYAEIEEIEE 299
|
|ESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYVHPRPGQDGAQAGVDGTV 165
|||:||||:|
|L-----PAPPTIAL-----SPALRQPCQARVLAT-----LRGQVLLSRAVGTAV 343
|
|ARINSSPLRYN-----RQIGEFIVTRAGLYLYL 199
|||:||||:|
|RRQLSREKRTFTVLAVVIGVFLVCWPFPPFFESY 381
|
|ELIMINARY; PRT; 532 AA.
|
|EMBLrel. 24, Created)
|EMBLrel. 24, Last sequence update)
|EMBLrel. 25, Last annotation update)
|date.
|vermitilis.
|nobacteria; Actinobacteridae; Actinomycetales;
|e; Streptomycetaceae; Streptomycetes.
|103;
|N.A.
| / ATCC 31267 / NCIMB 12804 / NRRL 8165;
|03; PubMed:11572948;
|la H., Ishikawa J., Hanamoto A., Takahashi C.,
|akahashi Y., Horikawa H., Nakazawa H., Osonoe T.,
|uba T., Sakaki Y., Hattori M.;
|ce of an industrial microorganism Streptomycetes
|educing the ability of producing secondary
|ad. Sci. U.S.A. 98:12215-12220(2001).
|N.A.
| / ATCC 31267 / NCIMB 12804 / NRRL 8165;
|06; PubMed:12692562;
|kawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba T.,
|tori M., Omura S.;
|me sequence and comparative analysis of the industrial
|Streptomycetes avermitilis."
|1. 21:526-531(2003).
|; BAC711174.1;
|; F-metalloendopeptidase activity; IEA.
|; Proteolysis and peptidolysis; IEA.
|02886; Peptidase_M37.
|; Peptidase_M37; I.
|ome.
|; AA; 55998 MW; 6E3F1CEC61E5A738 CRC64;
|7.2%; Score 103.5; DB 16; Length 532;
|urity 23.4%; Pred. No.1.4;
|nservative 22; Mismatches 107; Indels 103; Gaps 14
|
|SLGSRDGGAVRQAQPPAPMAARRSRRRRGRRGPGTALLVPLALGLALACGLL 73
|
|-----GSR-----AASTASRRRIIPARSALLTVAVPSACVGVNA-----GTA 316
|
|SLGSRASLSAQEPAQ-----BELVAEED-----QDPS 104
|
|3VGSDDSKAHTTASADAIAVKPSAANKLDTQIESLSAGADDFADRASRTQERI 376
|
|QTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYVHPRPGQDGAQ-----158
|QAAERKQKAAA---EAARKERLRPKFALPVAQKLSAVY-----GOAGINMWSVH 427

```

QY	159	-----AGVDGTIV-SGWEERARINSSSPLRYNROI GFIVTRAGLYLII
DB	428	TGIDFPVSYGTTVLAAATDGTVRQTWNSAYGNMA-----IVTAKDGTETWY
QY	204	FDECKAVYLKLDLLVDGVLAIRCLEEFSAATAASSLGPOLRLCQVSGILLARPQSLSL
DB	475	LSTYKVASGTTVKAGDPPI-----AFSCNCSNTGPHLH-----FEVYPAGGSJ
QY	264	LPW 266
DB	521	LPW 523
RESULT 10		
Q9KI66		
ID	Q9KY766	PRELIMINARY; PRT; 565 AA.
OS	Q9KY66;	
DT	01-OCT-2000 (TrEMBLrel. 15, Created)	
DT	01-OCT-2000 (TrEMBLrel. 15, Last sequence update)	
DE	01-JUN-2003 (TrEMBLrel. 24, Last annotation update)	
DE	Putative peptidase	
GN	SCO4798 OR SCD63A.09C.	
OS	Streptomyces coelicolor.	
OC	Bacteria; Actinobacteriae; Actinobacteridae; Actinomycetales;	
OC	Streptomycinae; Streptomycetaceae; Streptomyces.	
NCBI_TaxID=1902;		
RN	[1]	
RN	SEQUENCE FROM N.A.	
RC	STRAIN=A3(2);	
RA	Brown S.P., Harris D.;	
RL	Submitted (MAY-2000) to the EMBL/GenBank/DDBJ databases.	
RN	[2]	
RN	SEQUENCE FROM N.A.	
RC	STRAIN=A3(2);	
RA	Cerdeno A.M., Parkhill J., Barrell B.G., Rajandream M.A.;	
RL	Submitted (MAY-2000) to the EMBL/GenBank/DDBJ databases.	
RN	[3]	
RN	SEQUENCE FROM N.A.	
RC	STRAIN=A3(2);	
RX	MEDLINE=970000351; PubMed=8843436;	
RC	Redenbach M., Kieser H.M., Denapaitte D., Eichner A., Cullum J.,	
RA	Kinashi H., Hopwood D.A.;	
RT	"A set of ordered cosmids and a detailed genetic and physical map	
RT	of the 8 Mb Streptomyces coelicolor A3(2) chromosome."	
RL	Mol. Microbiol. 21:77-96(1996).	
RN	[4]	
RN	SEQUENCE FROM N.A.	
RC	STRAIN=A3(2) / M145;	
RC	MEDLINE=21996410; PubMed=12000953;	
RX	Bentley S.D., Chater K.F., Cerdeno-Tarraga A.-M., Challis G.L.,	
RA	Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,	
RA	Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M	
RA	Cronin A., Fraser A., Goble A., Hidalgo J., Hornaby T., Howarth S	
RA	Huang C.-H., Kieser T., Larke L., Murphy L., Oliver K., O'Neill S.	
RA	Rabinowitsch E., Rajandream M.A., Rutherford K., Rutter S.,	
RA	Seeger K., Saunders D., Sharp S., Squares R., Squares S., Taylor I,	
RA	Warren T., Wietzorek A., Woodward J., Barrell B.G., Parkhill J.,	
RA	Hopwood D.A.;	
RT	"Complete genome sequence of the model actinomycete Streptomyces	
RT	coelicolor A3(2)."	
RL	Nature 417:141-147(2002).	
DR	EMBL; AL2939121; CAB92661.1; "	
DR	GO; GO:0004222; F:metallopeptidase activity; IEA.	
DR	GO; GO:0006508; P:proteolysis and peptidolysis; IEA.	
DR	InterPro; IPR002886; Peptidase_M37.	
DR	Pfam; PF01551; Peptidase_M37; I.	
XO	Complete proteome.	
XO	SEQUENCE 565 AA.	
SO	SEQUENCE 565 AA.	

Query Match 7.1%; Score 103; DB 16; Length 565;
Best Local Similarity 21.2%; Pred. No. 1.7;
Matches 70; Conservative 33; Mismatches 99; Indels 128; G.

FEISA---RLPLPRSLGRSDGAVRQAQPPAPMAARRSQRRRGRRPGGTALL-- 56
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DOEATPDARIPVARA-GSRAGARSRRQP-----AKRS-----ALLTI 331
ALGI---GLALACLGILLAWVLSGRASLSAOEPAAEE----- 94
ACWMSVAGIAAASVGSLTG--DEGTTTASADPDCNAEAAPVKPSAANKLDTOLT 389
EED-----QDPSELNPQTEESQDPAPFLNLRVPRSPKPKRTRARRAIAAHY 146
GADDFADRSTQERIDLKAQDAEKRAAQEAARKERLRPKFALPVKHGLSAYV 449
RPGDQG-----AAQGVDTGTVSGWEAREINSSPIRYNRQICE 187
--GAQGINWWSHSGIDFPVLQGTITWMAATDGTVR-----TFINSAYGN 491
PA--GLYLXLCQVH-----FDGEKAVILKLDLLVDGVLAIRCLSEFSATAAS 236
TAKDGTETWTCHLSSVQVPSTTVKAGDAI-----AYSQDSGN 533
QLRLCQVSGLLAIRPGSSLRIRTL PW 266
HLH-----FEVRPAGGSSIDL PW 556

RELIMINARY; PRT; 955 AA.

TREMBRel. 24, Created)
TREMBRel. 24, Last sequence update)
TREMBRel. 24, Last annotation update)
protein OSUNB006008.10.
.10.
(japonica cultivar-group).
ridiplantae; Streptophyta; Embryophyta; Tracheophyta;
; Magnoliophyta; Liliopsida; Poales; Poaceae;
; Oryzaeae; Oryza.
947;

[N.A.
pponbare;
uan Q., Ouyang S., Liu J., Gansberger K., Jones K.M.,
L., Tstrin T., Kim M.M., Bera J.J., Jin S.S.,
Tallon L.J., Koo H., Zismann V., Hsiao J., Blunt S.,
Riedmuller S.B., Utterback T.T., Feildlyum T.V.,
as B.J., Suh B.B., Peterson J.J., Quackenbush J.,
zberg S.L., Fraser C.M.;
chromosome 3 BAC OJGNB0006008 genomic sequence." ;
Y-2002) to the EMBL/GenBank/DBJ databases.

[N.A.
pponbare;

R-2003) to the EMBL/GenBank/DBJ databases.
'6; AAO66523.1; --
Protein.
15 AA; 105582 MW; E44E88COFF71CC9C CRC64;

7.0%; Score 101.5; DB 10; Length 955;
arity 25.7%; Pred. No. 4.5;
Conservative 23; Mismatches 46; Indels 35; Gaps

ARRLPLPRSLGRSDGAVRQAQPPAPMAARRSQRR--GRRGPGTALLVPLAIGUG 64
|||
:::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
QNARPLP-----GSLMRAPPPPPTAEAPRQLPGAASAPATNTAA---- 180

ALGILLAWVLSGRASLSACEPAQEELVAEEDQPSLENPQTESODPAPFL---- 120
:::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
EPWILKGLVKPMSPQSIGPNRPSONE---DKDESE---EEEEEGFVIPDRA 232
-----NRLVPRRSAP 131

06:25:24 2004

us-09-245-198a-4.rspt

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EW e/threonine-protein kinase; Complete proteome.
EQ 3 AA; 92763 MW; CC4DA95AFAFE2407 CRC64;
CY 7.0%; Score 100.5; DB 16; Length 893;
  arity 22.7%; Pred.No. 5.1;
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  RLPIVR----RGELVRRRRARLPATTPHAKHAKDGRRPR-----PKRLGRNLVL 660
CY LLLAVVSL-----GSRA-----SLSAQEPAQEELVAEEDQDPSELN 107
  :|||:
  :|||:
  :|||:
DE NLAAAVAYAVMFMKASPNSDSGSKSDGRTGAAGGVSPAPEQSGGGSEPRPDQTS 720
CY ESQDPAPFLNRLIVRRPSAPKPKRTRARRAIAHYEVHPRGQDCAQAGVDGTVSG 167
  :|||:
  :|||:
  :|||:
DE KS-----PSGSSGSTREQTDPDVAQGFLLKDP--EGFRVAV---ATG 763
CY RINSSPLRYNKRQIGETIV 190
  :|||:
DE PKNGRGRIVYSHGDFELIV 786
CY
DE
SE April 7, 2004, 17:46:45
  secs
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GenCore version 5.1.1.6
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a search, using sw model

il 7, 2004, 17:38:07 ; Search time 11.7171 seconds
(without alignments)
1262.081 Million cell updates/sec

09-245-198A-4

SLDFFSARLPLRSLG.....PWAHLKAAPFLTYFGLFQV 284

SUM62

op 10.0 , Gapext 0.5

681 seqs, 52070155 residues

s satisfying chosen parameters: 141681

th: 0

th: 2000000000

nimum Match 0%

ximum Match 100%

sting first 45 summaries

issProt_42.*

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
d by analysis of the total score distribution.

SUMMARIES

ch	Length	DB	ID	Description
1.8	249	1	TN12 HUMAN	O43508 homo sapien
1.6	225	1	TN12 MOUSE	O54907 mus musculus
1.6	272	1	TNF5 CHICK	Q918d8 gallus gall
1.5	260	1	TNF5 CANFA	O97626 canis famil
1.4	254	1	TNF9 HUMAN	P41273 homo sapien
1.9	952	1	HDA7 HUMAN	O8wni4 homo sapien
1.7	441	1	CG22 ANIMA	P34801 antirrhinum
1.6	201	1	TNFB MACEU	Q9xt48 macropus eu
1.5	280	1	TNF6 MACMU	Q9myl6 macaca mula
1.5	690	1	RHO MICLU	P52154 micrococci
1.4	310	1	Y497 MYCTU	Q11162 mycobacteri
1.4	240	1	TN14 HUMAN	O43557 homo sapien
1.4	280	1	TNFC CERTO	Q9bdl1 cercocebus
1.4	902	1	NFC4 HUMAN	Q14934 homo sapien
1.3	760	1	MLH1 MOUSE	Q9jks1 mus musculus
1.3	814	1	CADF HUMAN	P55291 homo sapien
1.2	707	1	JIPI MOUSE	O9wv19 mus musculus
1.2	280	1	MDCB KLEPN	P71422 klebsiella
1.2	316	1	TN11 MOUSE	O35235 m tumor nec
1.1	278	1	TNFB RAT	P36940 rattus norv
1.1	281	1	TNF6 HUMAN	P48023 homo sapien
1.1	422	1	GF11 HUMAN	Q99684 homo sapien
1.1	574	1	SEN3 HUMAN	O9b414 homo sapien
1.0	197	1	TNFB RABIT	P10154 cryptolagus
1.0	204	1	TNFB BOVIN	Q06600 bos taurus
1.0	291	1	TN10 MOUSE	P50592 mus musculus
1.0	250	1	TNFC MACEU	Q9xt47 macropus eu
1.0	139	1	YQFB BACSU	P54467 bacillus eu
1.0	205	1	TNFB MARMO	Q9jmo9 marmota mon
1.0	241	1	TN13 MOUSE	Q9d777 mus musculus
1.0	777	1	METE CAUCR	Q9aaw1 caulobacter
1.0	933	1	VGLB_HSV1	O44463 herpesvirus
1.0	928	1	VGLB_HSVBP	P17471 bovine herp

RESULT 1	TN12 HUMAN	STANDARD;	PRT;	249 AA.
34	85.5	5.9	932	1 VGLB_HSVBP
35	85	5.9	372	1 LMXB_MOUSE
36	85	5.9	379	1 LMXB_HUMAN
37	84.5	5.9	401	1 AROC_MYCTU
38	84	5.8	284	1 TLX2_HUMAN
39	84	5.8	310	1 TNFC_MARMO
40	84	5.8	575	1 MIS_PIG
41	84	5.8	703	1 ZM15_HUMAN
42	84	5.8	825	1 ICP0_HSV2H
43	83.5	5.8	416	1 RAGE_BOVIN
44	83.5	5.8	505	1 TUB_MOUSE
45	83.5	5.8	545	1 RTN2_HUMAN

ALIGNMENTS

AC O43508: O8WUZ7;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DE Tumor necrosis factor ligand superfamily member 12 (TNF-related w
 DE inducer of apoptosis) (TWEAK) (APO3 ligand).
 GN TNFSF12 OR APO3L OR DR3LG.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A., AND N-TERMINUS OF SOLUBLE FORM.
 RC TISSUE=Fetal liver, and Tonsil;
 RX MEDLINE=98070415; PubMed=9405449;
 RA Chicheportiche Y., Bourdon P.R., Xu H., Hsu Y.-M., Scott H.,
 RA Hession C., Garcia I., Browning J.I.;
 RA "TWEAK, a new secreted ligand in the tumor necrosis factor family
 RT weakly induces apoptosis.";
 RL J. Biol. Chem. 272:32401-32410(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Fetal kidney;
 RX MEDLINE=9828355; PubMed=9560343;
 RA Marsters S.A., Sheridan J.P., Pitti R.M., Brush J., Goddard A.,
 RA Ashkenazi A.;
 RA "Identification of a ligand for the death-domain-containing recep
 RT Apo3.";
 RL Curr. Biol. 8:525-528(1998).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Tonsil;
 RX MEDLINE=23388257; PubMed=12477932;
 RA Strausberg R.L., Feigold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalish D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).

P12640 bovir
 O88609 mus t
 O60663 homo
 P95013 mycot
 O43763 homo
 Q9jmi0 marmc
 P79295 sus e
 Q9h091 homo
 P28284 herpe
 Q28173 bos t
 P50586 mus n
 O75298 homo

061: PubMed=10085077;
 ang Y.C., Lund J.K., Chen Y.-W., Leal J.A., Wiley S.R.;
 s angiogenesis and proliferation of endothelial cells.";
 .. 274:8455-8459(1999).
 Binds to FN14 and possibly also to TNFRSF12/AP03. Weak
 f apoptosis in some cell types. Mediates NF-KappaB
 n. May promote angiogenesis and the proliferation of
 al cells.
 Homotrimer (Potential).
 AR LOCATION: Type II membrane protein and secreted.
 ECIFICITY: Highly expressed in adult heart, pancreas,
 muscle, brain, colon, small intestine, lung, ovary,
 spleen, lymph node, appendix and peripheral blood
 es. Low expression in kidney, testis, liver, placenta,
 d bone marrow. Also detected in fetal kidney, liver,
 brain.
 soluble form derives from the membrane form
 lytic processing.
 Y: Belongs to the tumor necrosis factor family.
 Ref 3 sequence differs from that shown due to a
 t in position 125.

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 ail to license@isb-sib.ch).

 9; AAC51923.1; -;
 2; AAC39724.1; -;
 7; AAH19047.1; ALT_FRAME.
 1927; TNFSF12.

 7; C: integral to plasma membrane; TAS.
 2; P: receptor binding; TAS.
 7; P: induction of apoptosis; TAS.
 5; P: signal transduction; TAS.
 006052; TNF family.
 008983; TNF_like.
 7; TNF; 1.
 7; TNF; 1.
 251; TNF 1; FALSE_NEG.
 049; TNF2; 1.
 (ogenesis; Apoptosis; Transmembrane; Glycoprotein;
 1 249 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
 MEMBER 12, MEMBRANE FORM.
 94 249 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
 MEMBER 12, SECRETED FORM.
 1 21 CYTOPLASMIC (POTENTIAL).
 22 42 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
 (POTENTIAL).
 43 249 EXTRACELLULAR (POTENTIAL).
 93 94 CLEAVAGE.
 39 139 N-LINKED (GLCNAC...).
 19 AA; 27216 MW; E660843361C28EBA CRC64;
 87.8%; Score 1268; DB 1; Length 249;
 larity 100.0%; Pred. No. 4e-92;
 Conservative 0; Mismatches 0; Indels 0; Gaps 0;

 RSQRGRGPGTALLVPLALGLALACUGLLAVSLGSRASLSAQEPAQEL 95
 |||||
 RSQRGRGPGTALLVPLALGLALACUGLLAVSLGSRASLSAQEPAQEL 60
 |||||
 EDDPSLNPQTESQDPAPFLNRLVPRRSAPGKTRARRAIAAHYVHPRPGD 155
 |||||
 EDDPSLNPQTESQDPAPFLNRLVPRRSAPGKTRARRAIAAHYVHPRPGD 120
 |||||

QY 156 GAQAGVDGTSGWBEARINSSPLRYNRQIGFIVTRAGLYLYLYCOVHFDEGKAVY
 Db 121 GAQAGVDGTSGWBEARINSSPLRYNRQIGFIVTRAGLYLYLYCOVHFDEGKAVY
 QY 216 LLDVGVIALRLCLERFSATAASSLGPQLRLCOVSGLLALRPGSSLRIRTLPAHLKA
 Db 181 LLDVGVIALRLCLERFSATAASSLGPQLRLCOVSGLLALRPGSSLRIRTLPAHLKA
 QY 276 TYFGLFQVH 284
 Db 241 TYFGLFQVH 249

 RESULT 2
 TN12 MOUSE
 ID TN12 MOUSE STANDARD; PRT; 225 AA.
 AC 054907; OSCPT2;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, last sequence update)
 DT 28-FEB-2003 (Rel. 41, last annotation update)
 DE Tumor necrosis factor ligand superfamily member 12 (TNF-related
 DE inducer of apoptosis) (TWBAK) (Fragment).
 GN TNFSF12.
 OS Mus musculus (Mouse).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus
 CC NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-Peritoneal macrophage;
 RX MEDLINE=98070415; PubMed=9405449;
 RA Chicheportiche Y., Bourdon P.R., Xu H., Hsu Y.-M., Scott H.,
 RA Hession C., Garcia I., Browning J.L.;
 RT "TWEAK, a new secreted ligand in the tumor necrosis factor family;
 RT weakly induces apoptosis".
 RL J. Biol. Chem. 272:32401-32410(1997).
 RN [2]
 RP SEQUENCE OF 93-225 FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Retina;
 RX MEDLINE=21085660; PubMed=11217851;
 RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.
 RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.
 RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamataka I.
 RA Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito K.
 RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
 RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
 RA Kuehl P., Lewis S., Mateuo Y., Nikaudo I., Pesole G., Quackenbush
 RA Schriml L.M., Stauber E., Suzuki R., Tomita M., Wagner L., Washi
 RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
 RA Blake J., Boileau D., Bojunga N., Carninci P., de Bonaldo M.F.,
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
 RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.I.
 RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
 RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilmi
 RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohsaki S.
 RA Hayashizaki Y.;
 RT "Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 409:685-690(2001).
 CC -!- FUNCTION: Binds to FN14 and possibly also to TNFSF12/AP03. I
 inducer of apoptosis in some cell types. Promotes angiogenes
 the proliferation of endothelial cells. Mediates NF-KappaB
 activation (By similarity).
 CC SUBUNIT: Homotrimer (Potential).
 CC -!- SUBCELLULAR LOCATION: Type II membrane protein and secreted
 similarity).
 CC -!- TISSUE SPECIFICITY: Widely expressed.
 CC -!- PTM: The soluble form is produced from the membrane form by
 proteolytic processing (By similarity).
 CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a col.

STANDARD; PRT; 260 AA.

Rel. 40, Created)
 Rel. 40, Last sequence update)
 Rel. 41, Last annotation update)
 s factor ligand superfamily member 5 (CD40 ligand).
 OLIG OR CD40L.

ris (Dog).
 tazia; Chordata; Craniata; Vertebrata; Euteleostomi;
 heria; Carnivora; Fissipedia; Canidae; Canis.
 15;

N.A.
 illett B.J.;
 perities of Canine CD40L";
 G-1998) to the EMBL/GenBank/DBJ databases.
 Cytokine that binds to TNFRSF5. Mediates B-cell
 tion in the absence of co-stimulus as well as iGe
 n in the presence of IL-4. Involved in immunoglobulin
 tching (By similarity).
 Homotrimer (By similarity).
 AR LOCATION: Type II membrane protein. Also exists as an
 ular soluble form (By similarity).
 soluble form derives from the membrane form by
 ic processing (By similarity).
 y: Belongs to the tumor necrosis factor family.

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 ail to license@isb-sib.ch).

1; AAD04375.1; -.
 1ALY.
 1; C: integral to membrane; ISS.
 4; F: CD40 receptor binding; ISS.
 10; P: B-cell proliferation; ISS.
 4; P: inflammatory response; ISS.
 9; P: leukocyte cell adhesion; ISS.
 8; P: platelet activation; ISS.
 003263; TNF 5.
 006052; TNF family.
 008983; TNF-like.
 003636; TNF_subf.
 ; TNF; 1.
 02; CD40LIGAND.
 1600; TNF 5; 1.
 012; TNF_subf; 1.
 17; TNF; 1.
 251; TNF 1; 1.
 049; TNF 2; 1.
 usmembrane; Glycoprotein; Signal-anchor.

1 260
 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY

12 260
 MEMBER 5, MEMBRANE FORM.

12 260
 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY

1 22
 MEMBER 5, SOLUBLE FORM (BY SIMILARITY).

23 46
 CYTOPLASMIC (POTENTIAL).

47 260
 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)

111 112
 (POTENTIAL).

177 217
 EXTRACELLULAR (POTENTIAL).

239 239
 CLEAVAGE (BY SIMILARITY).

50 AA; 28688 MW; 604F69A19E98EB70 CRC64;
 7.5%; Score 108.5; DB 1; Length 260;
 Larity 25.5%; Pred.No. 0.16;
 Conservative 23; Mismatches 69; Indels 63; Gaps 8;

103 EMKEENIAMQKGDQDPRIAAHVISEASSNPASVL-----RWAPKGYTISNNI
 146 YEYHPRGQDGAQAGVDGTSGWEEARINSSPLRYNKGIFVTRAGLYLYCC
 155 -----ENGKQ-----LAVKQGLYYVYAC
 206 EGKAVYLKLDLLVDGVLALRCLEFSAT-----AASLGFQLRLCQVS-----GLI
 178 SNRAASQAPF-----VASLCLHSPSGTERVLLRAASSRGSSKPCGQQSILHGGVF
 256 GSSLRIRTLPAWHLKAAPFLTYFGLFQV 283
 233 GASVFVNVTDPQSVSHGTGTFTSGLLKL 260

RESULT 5
 TNF9 HUMAN
 ID TNF9_HUMAN STANDARD; PRT; 254 AA.
 AC P41273;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Tumor necrosis factor ligand superfamily member 9 (4-LBB ligand)
 DE 1LBI.
 GN TNFSF9.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_Taxid=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=94374434; PubMed=8088337;
 RA Alderson M.R., Smith C.A., Tough T.W., Davis-Smith T., Armitage F.
 RA Falk B., Roux E., Baker E., Sutherland G.R., Din W.S., Goodwin R.
 RT "Molecular and biological characterization of human 4-LBB and its
 ligand.";
 RL Eur. J. Immunol. 24:2219-2227(1994).
 CC -!- FUNCTION: Cytokine that binds to TNFSF9. Induces the
 proliferation of activated peripheral blood T cells. May have
 role in activation-induced cell death (AICD). May play a role
 cognate interactions between T cells and B cells/macrophages.
 CC -!- SUBUNIT: Homotrimer (Potential).
 CC -!- SUBCELLULAR LOCATION: Type II membrane protein.
 CC -!- TISSUE SPECIFICITY: EXPRESSED IN BRAIN, PLACENTA, LUNG, SKIN
 MUSCLE AND KIDNEY.
 CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.

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 modified and this statement is not removed. Usage by and for
 entities requires a license agreement (See <http://www.isb-sib.ch/>
 or send an email to license@isb-sib.ch).

EMBL; U03398; AAA53134.1; -.
 PIR; I38427; I38427.
 Genew; HGNC:11939; TNFSF9.
 MIM; 606182; -.
 GO; GO:0005102; F:receptor binding; TAS.
 GO; GO:0006915; P:apoptosis; TAS.
 GO; GO:0008283; P:cell proliferation; TAS.
 GO; GO:0007267; P:cell-cell signaling; TAS.
 GO; GO:0007165; P:signal transduction; TAS.
 InterPro; IPR006052; TNF family.
 InterPro; IPR008983; TNF-like.
 Pfam; PF00229; TNF; 1.
 SMART; SM00207; TNF; 1.
 PROSITE; PS00251; TNF 1; 1.
 PROSITE; PS00049; TNF 2; 1.
 Cytokine; Transmembrane; Glycoprotein; Signal-anchor; Polymorphi;
 DOMAIN 1 28
 CYTOPLASMIC (POTENTIAL).

9 49 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
(POTENTIAL).
0 254 EXTRACELLULAR (POTENTIAL).
5 41 POLY-LEU.
7 17 P-> A (in dbSNP:442511).
AA: 26624 MW; 827551F34563E508 CRC64;
7.4%; Score 106.5; DB 1; Length 254;
rity 26.5%; Pred.No.0.22; 93; Indels 81; Gaps 13;
nservative 31; Mismatches 93; Indels 81; Gaps 13;
AARSRRRRRRGGPGTALLVPLALGLALACGLGLLVVSL-GSRASL-SAQE 89
-----RACRVLPL-WALVAGLLLLLLAAACAVLACPAWVGARSPGSAAS 67
LVAEEDQDPPELNPQTESQDPAPFLNRLVPRPSAPKGRKTRARRAIAHYEVH 149
-----GPESLP-----DDPAGLLDL-----RQGMFAQLVAQNVL- 102
DGAQAGVDGTVSGWEE---ARINSSPLRYNRQIGFIVTRAGLYLYLCOWHFE 206
-----IDGPLSWYSDPLGAGVSLTGSLYKEDTKELVAKAGVYVFFQ----- 146
LKDLVLVDG-----VLALRCLEFSAATASSLPQRLCQVSG----- 249
LELRVVAGEGSGSVSLALHLQPLRSAGAALALTVDLPPASSEARNASFGQ 201
RPGSSLIRTLPLWAHLKAAPFL-----TYFGLFQV 283
SAGQRLGVHLHTEARHAWQLTGATVGLFRV 240
STANDARD; PRT; 952 AA.
; Q96K01; Q9BR73; Q9H7L0; Q9NW41; Q9NWA9; Q9NYK9;
el. 42, Created)
el. 42, Last sequence update)
el. 43, Last annotation update)
ylase 7a (HD7a).
7.
Human).
azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Catarrhini; Hominidae; Homo.
6;
N.A. (ISOFORM 1).
l carcinoma;
W., Verdin E., Walsh M.J.;
II HDAC is associated with the transcriptional
pressor CCAT displacement protein.";
-2000) to the EMBL/GenBank/DBJ databases.
N.A. (ISOFORM 2), SEQUENCE OF 220-952 FROM N.A. (ISOFORM
CE OF 651-952 FROM N.A.
Placenta, Spleen, and Teratocarcinoma;
T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,
Nagai K., Sugano S., Shiratori A., Sudo H.,
Hosokiri T., Kaku Y., Kodaira H., Kondo H., Sugawara M.,
Chiba Y., Ishida S., Murakawa K., Ono Y., Takiguchi S.,
Imura K., Murakami K., Ishii S., Kawai Y., Saito K.,
Ikanatsu A., Nakamura Y., Nagahari K., Masubo Y.,
wayanagi T.;
NA sequencing project.";
i-2003) to the EMBL/GenBank/DBJ databases.
N.A. (ISOFORM 3).
histone deacetylase 7A (HDAC7A), transcript variant 3.";
i-2003) to the EMBL/GenBank/DBJ databases.

RP SEQUENCE FROM N.A.
RA Worley K.C.;
RL Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.
RN [5]
RP SEQUENCE OF 75-952 FROM N.A. (ISOFORM 1).
RC TISSUE=Uterus;
RA Kohrer K., Beyer A., Mewes H.-W., Gassenhuber J., Wiemann S.;
RL Submitted (SEP-1999) to the EMBL/GenBank/DBJ databases.
RN [6]
RP SEQUENCE OF 241-952 FROM N.A. (ISOFORM 1).
RC TISSUE=B-cell, and Colon;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.,
RA Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.C.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmitz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalusz D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length t
and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [7]
RP SUBCELLULAR LOCATION, AND INTERACTION WITH EDNRA.
RX MEDLINE=21264398; PubMed=11262386;
RA Lee H.-J., Chun M., Kandroz K.V.;
RT "Tip60 and HDAC7 interact with the endothelin receptor a and may b
involved in downstream signaling.";
RL J. Biol. Chem. 276:16597-16600(2001).
RN [8]
RP INTERACTION WITH HDAC3.
RX MEDLINE=21443773; PubMed=11466315;
RA Fischle W., Dequiedt F., Fillion M., Hendzel M.J., Voelter W.,
RA Verdin E.;
RT "Human HDAC7 histone deacetylase activity is associated with HDAC3
vivo.";
RL J. Biol. Chem. 276:35826-35835(2001).
RN [9]
RP FUNCTION.
RX MEDLINE=22224741; PubMed=12239305;
RA Bryant H., Farrell P.J.;
RT "Signal transduction and transcription factor modification during
reactivation of Epstein-Barr virus from latency.";
RL J. Virol. 76:10290-10298(2002).
RN [10]
RP INTERACTION WITH HTATIP.
RX MEDLINE=22538515; PubMed=12551922;
RA Xiao H., Chung J., Kao H.-Y., Yang Y.-C.;
RT "Tip60 is a co-repressor for STAT3.";
RL J. Biol. Chem. 278:11197-11204(2003).
CC -!- FUNCTION: Responsible for the deacetylation of lysine residues
the N-terminal part of the core histones (H2A, H2B, H3 and H4)
Histone deacetylation gives a tag for epigenetic repression at
plays an important role in transcriptional regulation, cell cy
progression and developmental events. Histone deacetylases act
the formation of large multiprotein complexes. Involved in mus
maturation by repressing transcription of myocyte enhancer fac
such as MEF2A, MEF2B and MEF2C. During muscle differentiation,
shuttles into the cytoplasm, allowing the expression of myocyte
enhancer factors (By similarity). May be involved in Epstein-B
virus (EBV) latency, possibly by repressing the viral BZLF1 ge
-!- SUBUNIT: Interacts with HDAC1, HDAC2, HDAC3, HDAC5, NCO
NCOR2, SIN3A, SIN3B, RBBP4, RBBP7, MTA1L1, SAP30 and MED3.

with the 14-3-3 protein YWAE, MEF2A, MEF2B and MEF2C activity). Interacts with HTATIP and EDNRA. AR LOCATION: Nuclear and cytoplasmic. In the nucleus, it has with distinct subnuclear dot-like structures. Shuttles in nucleus and the cytoplasm. Treatment with EDN1 results in the nucleus to the perinuclear region. The cytoplasm depends on the interaction with the 14-3-3 YWAE and may be due to its phosphorylation. VE PRODUCTS:

alternative splicing; Named isoforms=4;

8WU14-1; Sequence=Displayed;

8WU14-2; Sequence=VSP_007429, VSP_007431;

experimental confirmation available;

8WU14-4; Sequence=VSP_008772;

8WU14-3; Sequence=VSP_007430; experimental confirmation available; he nuclear export sequence mediates the shuttling between the nucleus and the cytoplasm (By similarity).

be phosphorylated by Cdk1 (By similarity).

FOUS: Its activity is inhibited by Trichostatin A (TSA), histone deacetylase inhibitor (By similarity).

Y: Belongs to the histone deacetylase family. Subfamily

Ref.1 sequence differs from that shown due to a

t in position 877.

Ref.2 (BAC56929) sequence differs from that shown due to

etention.

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3; AAF63491.1; ALT_FRAME.

2; BAA91474.1; ALT_INIT.

0; BAA91545.1; ALT_INIT.

9; BAB15759.1; ALT_INIT.

1; BAB55363.1; ALT_INIT.

8; BAC56929.1; ALT_SEQ.

7; AAP84704.1; -.

6; -; NOT_ANNOTATED_CDS.

5; CAB55935.1; -.

3; AAH06453.1; ALT_INIT.

5; AAH20505.1; ALT_INIT.

4067; HDAC7A.

7; Cytoplasm; TAS.

8; Histone deacetylase complex; TAS.

4; C:nucleus; TAS.

7; F:histone deacetylase activity; TAS.

6; F:specific transcriptional repressor activity; TAS.

4; F:transcription factor binding; TAS.

13; P:B-cell differentiation; TAS.

13; P:chromatin modification; TAS.

13; P:inflammatory response; TAS.

13; P:negative regulation of myogenesis; TAS.

13; P:neurogenesis; TAS.

14; P:regulation of cell cycle; TAS.

1000286; His_deacetylase.

1; Hist_deacetyl; 1.

270; HDASUPER.

nuclear protein; Chromatin regulator;

1 regulation; Repressor; Repeat; Phosphorylation;

splicing.

1 269 TRANSCRIPTION REPRESSION 1 (BY

SIMILARITY).

FT	DOMAIN	218	546	TRANSCRIPTION REPRESSION 2 (BY SIMILARITY).
FT	DOMAIN	518	865	HISTONE DEACETYLASE.
FT	DOMAIN	918	952	NUCLEAR EXPORT (BY SIMILARITY).
FT	DOMAIN	1	98	INTERACTION WITH MEF2C (BY SIMILARITY).
FT	DOMAIN	49	149	INTERACTION WITH MEF2A (BY SIMILARITY).
FT	DOMAIN	877	952	INTERACTION WITH SIN3A (BY SIMILARITY).
FT	DOMAIN	197	203	POLY-SER.
FT	DOMAIN	368	373	POLY-PRO.
FT	ACT_SITE	670	670	BY SIMILARITY.
FT	VARSPLIC	1	472	Missing (in isoform 2).
FT	VARSPLIC	227	263	/FTid=VSP_007429.
FT	VARSPLIC	227	256	Missing (in isoform 3).
FT	VARSPLIC	227	256	/FTid=VSP_008772.
FT	VARSPLIC	227	256	Missing (in isoform 4).
FT	VARSPLIC	227	256	/FTid=VSP_007430.

Query Match 6.9%; Score 99; DB 1; Length 952;
Best Local Similarity 24.2%; Pred. No. 3.9;
Matches 46; Conservative 21; Mismatches 67; Indels 56; G

Qy	11	RRLLPRLSLGRDGGAVR	-----QAOPPAPMAAR-----RSQRR
Db	401	RRLLPRLSLGRDGGAVR	-----QAOPPAPMAAR-----RSQRR
Qy	48	-RGEPFTALLVPLALGLALACILGLLLAVVSLGRSLASLQAEPAEELVAEDQI	-----RAQSSPAAPASLQSAPEPASQARVLSSET
Db	461	PRGSTGTDTLLPLAQGGHRLS	-----RAQSSPAAPASLQSAPEPASQARVLSSET
Qy	107	NPQT	-----EESQDP-----
Db	516	LPFTTGLIYDVMUKHQCSDGNSRHPHAGRIQSIWSRLQERGLRSCCELRGR	-----APFLNLRVRRSPAPKGRK
Qy	140	RAIAAHVEVH	149
Db	576	ELQSVHSEH	585

RESULT 7

ID	CG22	ANTMA	STANDARD;	PRT;	441	AA.
AC	P34801;					
DT	01-FEB-1994	(Rel. 28, Created)				
DT	01-FEB-1994	(Rel. 28, Last sequence update)				
DT	16-OCT-2001	(Rel. 40, Last annotation update)				
DE	G2/mitotic-specific cyclin 2.					
OS	Antirrhinum majus (Garden snapdragon).					
OC	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;					
OC	Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; ast					
OC	lamiales; Lamiales; Antirrhinaceae; Antirrhineae; Antirrhinum.					
OX	NCBI_TaxID=4151;					
RN	[1]					
RP	SEQUENCE FROM N.A.					
EX	MEDLINE=94148008; PubMed=83113906;					
RA	Robert P.R., Coen E.S., Murphy G.J.P., Doonan J.H.;					
RT	"Patterns of cell division revealed by transcriptional regulation					
RT	genes during the cell cycle in plants."					
RL	EMBO J. 13:616-624(1994).					
CC	FUNCTION: Essential for the control of the cell cycle at the					
CC	(mitosis) transition. G2/M cyclins accumulate steadily during					
CC	and are abruptly destroyed at mitosis.					
CC	SUBUNIT: Interacts with the CDC2 and CDK2 protein kinases to					
CC	a serine/threonine kinase holoenzyme complex. The cyclin sub					
CC	imparts substrate specificity to the complex.					
CC	DEVELOPMENTAL STAGE: Accumulates steadily during G2 and is					
CC	abruptly destroyed at mitosis.					
CC	SIMILARITY: Belongs to the cyclin family. Cyclin AB subfamil					
CC	This SWISS-PROT entry is copyright. It is produced through a col					
CC	between the Swiss Institute of Bioinformatics and the EMBL out					
CC	the European Bioinformatics Institute. There are no restriction					
CC	use by non-profit institutions as long as its content is in					
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CAAS3729.1; -;
41710.
1VIN.
06670; Cyclin.
04367; Cyclin Cterm.
06671; Cyclin_N.
cyclin_C; 1.
cyclin_C; 1.
CYCLIN; 2.
92; CYCLINS; 1.
ycle; Cell division; Mitosis.
AA; 49205 MW, E6E4C037C98880A7 CRC64;

6.7%; Score 97; DB 1; Length 441;
rity 24.0%; Pred. No. 2.3;
nservative 44; Mismatches 104; Indels 74; Gaps 15;

QAPPAPMAARRSQR-----RGRGPGTALVPLALGLALACLGLL 73

KS-----MAVEKKRRALDIGNVTVRGKALPQVSRPIIRGF-----CAQLI 69

-----LAVSLGRASLS-----AQEPAQEELVAEEDQDPSELNPQTEESQ 114

AAENNNKSLAVNAGADGALPIKEAVARVPQKTKVSKQEIIIEISPDTEKKX 129

LNRLVRRRS-----APGRKTRARRAIAHVEVHPRPGQ-----DGAAGVDGTV 165

LEKEITGSKLKKKAPTTLTSTLTARSKAASV-VRTKPKQIVDIDAADVNDLAV 186

-----ARINSSPLRY-----NRQIGFIVTRAGLYLYLCVHFDP-----EGKAVYL 212

DMYFKYGAENDSRPHDYMDSQPEINEKK-----RAILIDLWLVQVHYKFEISPETLYL 244

NDGVILALRC-----LEEFSATAASSLGPQLRLCQVSGLLALRPGS 257

VDVYLASKTSRRELQLLGMSSMLIASKYEEIWAPEVNDLVCIISDGS 295

STANDARD; PRT; 201 AA.

rel. 40, Created)
rel. 40, last sequence update)
rel. 42, last annotation update)
pha precursor (LT-alpha) (TNF-beta) (Tumor necrosis
superfamily member 1).
OR TNFB.
lii (Tamar wallaby).
azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Diprotodontia; Macropodidae; Macropus.
5;

N.A.

48; PubMed=10826697;

Deane E.M.;

of lymphotoxin alpha (LT-alpha) from a marsupial,

lii;

9-403(2000).

Cytokine that in its homotrimeric form binds to

TNFR1, TNFRSF1B/TNFR and TNFRSF14/HVEM. In its

meric form with LTB binds to TNFRSF3/LTBR. Lymphotoxin is

by lymphocytes and cytotoxic for a wide range of tumor

nitro and in vivo.

homotrimer, and heterotrimer of either two LTB and one

its or (less prevalent) two LTA and one LTB subunits (By

).

AR LOCATION: Secreted (homotrimer) and membrane-

i (heterotrimers) (By similarity).

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CC the European Bioinformatics Institute. There are no restriction
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CC entities requires a license agreement (See <http://www.isb-sib.ch/>
CC or send an email to license@isb-sib.ch).

CC -----
CC EMBL; AF119336; AAD41773.1; -;
CC HSSP; P01374; LTNR.
CC InterPro; IPR006053; TNF abc.
CC InterPro; IPR006052; TNF family.
CC InterPro; IPR008983; TNF like.
CC InterPro; IPR003636; TNF_subf.
CC Pfam; PF00229; TNF; 1.
CC PRINTS; P01234; TNECROSISFCT.
CC ProDom; PD002012; TNF_subf; 1.
CC SMART; SM0207; TNF; 1.
CC PROSITE; PS00251; TNF 1; 1.
CC PROSITE; PS0049; TNF 2; 1.
CC CytoKine; Glycoprotein; Signal.
CC SIGNAL 1 27 BY SIMILARITY.

CC CHAIN 28 201 LYMPHOTOXIN-ALPHA.
CC CARBOHYD 93 93 N-LINKED (GLCNAC...) (POTENTIAL).
CC SEQUENCE 201 AA; 21536 MW; 8C4C371CB5091627 CRC64;

Query Match 6.6%; Score 95.5; DB 1; Length 201;
Best Local Similarity 23.2%; Pred. No. 1.2;
Matches 44; Conservative 27; Mismatches 84; Indels 35; G;

QY 107 NPQTESQDPAPFLNRLVRRSAPKGRKTRARRAI--AAHYEVHPRPGDGAQAG

Db 30 NPDNHHSSPAP-----PQTAQHLSQKSLKEITLKPAAHL-----VGDPSVQDS.

QY 165 VSGWEARINSSP-LRYNRQI--GEFIVTRAGLYLYLCVHFDEGKA-----V

Db 76 --W---RANTDHAFLRHGFSLSNNLSLVPTSGLYEYVSVQVWFSGASCSEITPTLL

QY 215 DLLVDG---VLALRCLEEFSAATAASSLGPQLRLCQVSGLLALRPGSSLRITLPMW

Db 130 EVLLFSSKYQVHPVLLSAQSVCSGTGQPMRSVYQCAVFLITQGRSLTYTDTGVS

QY 272 APFLTYFGLF 281

Db 190 SPSSVFFGAF 199

RESULT 9

TNF6_MACMU

ID TNF6_MACMU STANDARD; PRT; 280 AA.

AC Q9MYL6; Q9BDM5;

DT 28-FEB-2003 (Rel. 41, Created)

DT 28-FEB-2003 (Rel. 41, Last sequence update)

DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE Tumor necrosis factor ligand superfamily member 6 (FAS antigen li

DE (CD95L protein).

GN TNFSF6 OR FASL OR CD95L.

OS Macaca mulatta (Rhesus macaque),

OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey), an

OS Macaca nemestrina (Pig-tailed macaque).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheidae;

OC Cercopitheidae; Macaca.

OX NCBI_TaxID=9544, 9541, 9545;

RN [1]

SEQUENCE FROM N.A.

RC SPECIES=M.mulatta; TISSUE=Lymphocytes;

RX MEDLINE=21383618; PubMed=11491535;

RA Villinger F., Bostik P., Mayne A.E., King C.L., Genain C.P.,

RA Weiss W.R., Ansari A.A.;

RT "Cloning, sequencing, and homology analysis of nonhuman primate

Fas/Fas-ligand and co-stimulatory molecules.";

RL Immunogenetics 53:315-328(2001).

```

79  GLCLLVFFWNLVALVGLG--LGMQLFLHQLXEL-----AELRESTSQKHATA
122  RLVRPRRSAPKGRKTRARRAIAAHYEHVPRPQDGAQGVDTGSGWEA-RINSS
129  QIGHP---SPPEKKEQRK--VAHLTGKNSRSMPL-----WEDTYGIVL
181  YNRQIGEFIVTRAGLYLYLCQVHFDEGKA-----VYLKLD-----LLVDGV
175  YKK--GSLVINETGLYFVYSKYVF-RGOSCTNLPUSHKYVNRNSKYFODLVMMSGK
226  CLEEFSAATASSIGPOLRCQVSGLLALRPSSGLRIRTLPAWHLKAAPFLTYPGFL
232  CTTGQMAHSSYLGAVENTSDADLY-----VNVSELSLVNFEESQ--TFFGLY

RESULT 10
RHO_MICLU
ID   RHO_MICLU          STANDARD;             PRT;       690 AA.
AC   P52I54;
CD   01-OCT-1996 (Rel. 34, Created)
DT   01-NOV-1997 (Rel. 35, Last sequence update)
DE   28-FEB-2003 (Rel. 41, Last annotation update)
DE   Transcription termination factor rho.
GN   RHO.
OS   Micrococcus luteus (Micrococcus lysodeikticus).
OC   Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC   Micrococccineae; Micrococccaceae; Micrococccus.
OX   NCBI_TaxID=1270;
RN   [1]
RS   SEQUENCE FROM N.A., AND SEQUENCE OF 1-5 AND 289-297.
RC   STRAIN=EM;
RC   MEDLINE=96132802; PubMed=8557681;
RA   Nowatke W.L., Richardson J.P.;
RA   "Characterization of an unusual Rho factor from the high G + C gr:
RA   positive bacterium Micrococcus luteus.";
RA   J. Biol. Chem. 271:742-747(1996).
RN   [2]
RS   SEQUENCE OF 205-690 FROM N.A.
RC   STRAIN=EM;
RC   MEDLINE=94327472; PubMed=8051015;
RA   Opperman T., Richardson J.P.;
RA   "Phylogenetic analysis of sequences from diverse bacteria with
RA   homology to the Escherichia coli rho gene.";
RA   J. Bacteriol. 176:5033-5043(1994).
RN   [3]
RS   REVISION TO 500.
RC   STRAIN=EM;
RC   Nowatke W.L.;
RA   Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
CC   -!- FUNCTION: FACILITATES TRANSCRIPTION TERMINATION BY A MECHANISM
CC   THAT INVOLVES RHO BINDING TO THE NASCENT RNA, ACTIVATION OF I
CC   RNA-DEPENDENT ATPASE ACTIVITY, AND RELEASE OF THE MRNA FROM A
CC   RNA TEMPLATE. RNA-DEPENDENT NPASE WHICH UTILIZES ALL FOUR
CC   RIBONUCLEOSIDE TRIPHOSPHATES AS WELL AS DATP AS SUBSTRATES, I
CC   HAS A SIGNIFICANT LOWER ACTIVITY WITH CTP.
CC   -!- SUBUNIT: Homohexamer (By similarity).
CC   -!- SIMILARITY: Contains 1 RNA recognition motif (RRM) domain.
CC   -----
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CC   the European Bioinformatics Institute. There are no restriction
CC   use by non-profit institutions as long as its content is not
CC   modified and this statement is not removed. Usage by and for
CC   entities requires a license agreement (See http://www.isb-sib.ch/
CC   or send an email to license@isb-sib.ch).
CC   -----
CC   EMBL; L27277; AAB18671.1; -.
CC   HSSP; P03002; 1A63.
DR   DR
DR   InterPro; IPR003593; AAA_ATPase.
DR   DR
DR   InterPro; IPR000194; ATPase_a/bcentre.
DR   DR
DR   InterPro; IPR002059; Cold_shock.
DR   DR
DR   InterPro; IPR008994; Nucleic_acid_OB.
DR   DR
DR   InterPro; IPR004665; Term_rho.

```


[illegible]

heria; Primates; Catarrhini; Hominidae; Homo.
06;

(N.A. (ISOFORM 1)).

340; PubMed=9462508;
bner R., Montgomery R.I., Kochel K.D., Cheung T.C.,
en S., Murphy M., Eisenberg R.J., Cohen G.H., Spear P.G.,
member of the TNF superfamily, and lymphotoxin alpha are
erpesvirus entry mediator.";
-30(1998).

(N.A. (ISOFORM 1)), AND CHARACTERIZATION.

532; PubMed=9765287;
McDonnell P.C., Brigham-Burke M., Lyn S.D., Minton J.,
le K., Spanpanato J., Silverman C., Hensley P.,
Emery J.G., Deen K., Eichman C., Chabot-Fletcher M.,
ung P.R.;
entry mediator ligand (HVEM-L), a novel ligand for
mulates proliferation of T cells and inhibits HT29 cell
l. 273:27548-27556(1998).

(N.A. (ISOFORM 2)), AND PROCESSING.

948; PubMed=11673523;
Butrovich K.D., Houshmand P., Edwards W.R., Ware C.F.;
acterization of LIGHT reveals linkage to an immune
is on chromosome 1p13.3 and distinct isoforms generated
splicing or proteolysis.";
.67:5122-5128(2001).

(N.A.

1257; PubMed=12477932;

L., Feingold E.A., Grouse L.H., Derge J.G.,
Collins F.S., Wagner L., Shemen C.M., Schuler G.D.,
Zeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
Marusina K., Farmer A., Rubin G.M., Hong L.,
Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
Quellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
ton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
ladan A., Young A.C., Shevchenko Y., Bouffard G.G.,
Touchman J.W., Green E.D., Dickson M.C.,
Grimwood J., Schmutz J., Myers R.M.,
S.N., Krzywinski M.I., Skalska U., Smalios D.E.,
Schein J.E., Jones S.J.M., Marra M.A.;
and initial analysis of more than 15,000 full-length
cDNA sequences.";

acad. Sci. U.S.A. 99:16898-16903(2002).

Cytokine that binds to TNFRSF3/LTR. Binding to the
receptor TNFRSF6B modulates its effects. Activates NFkB,
as the proliferation of T cells, and inhibits growth of
xarcinoma HT-29. Acts as a receptor for Herpes simplex

Homotrimer.

AR LOCATION: Type II membrane protein and secreted
1); Cytoplasmic (isoform 2).

(VE PRODUCTS:

ernative splicing; Named isoforms=2;

X3557-1; Sequence-Displayed;

ynonyms=LIGHT delta-TW;

X43557-2; Sequence=VSP_006452;

PECIFICITY: PREDOMINANTLY EXPRESSED IN THE SPLEEN BUT ALSO
THE BRAIN. WEAKLY EXPRESSED IN PERIPHERAL LYMPHOID
AND IN HEART, PLACENTA, LIVER, LUNG, APPENDIX, AND KIDNEY,
(PRESSION SEEN IN FETAL TISSUES, ENDOCRINE GLANDS, OR

CC NONHEMATOPOIETIC TUMOR LINES.
CC -!- INDUCTION: UPREGULATED AFTER T-CELL ACTIVATION.
CC -!- PTM: N-glycosylated.
CC -!- PTM: The soluble form of isoform 1 derives from the membrane
CC by proteolytic processing.
CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
CC -!- CAUTION: Ref.4 sequence differs from that shown due to a
CC frameshift in position 178.

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CC use by non-profit institutions as long as its content is li
CC modified and this statement is not removed. Usage by and for
CC entities requires a license agreement (See <http://www.isb-sib.ch/>
CC or send an email to license@isb-sib.ch).

CC EMBL; AF036581; AAC39563.1; -;
CC EMBL; AF064090; AAC25169.1; -;
CC EMBL; AY028261; AAK26160.1; -;
CC EMBL; BC018058; AAH18058.1; ALT_FRAME.
CC HSSP; P01375; 4TSV.
CC Genew; HGNC:11930; TNFSF14.
CC MIM; 604520; -;

CC GO; GO:0005102; P:receptor binding; TAS.
CC GO; GO:0006917; P:induction of apoptosis; TAS.
CC GO; GO:0007165; P:signal transduction; TAS.

CC InterPro; IPR006053; TNF abc.
CC InterPro; IPR006052; TNF_family.
CC InterPro; IPR008983; TNF_like.
CC InterPro; IPR003636; TNF_subf.
CC Pfam; PF00229; TNF; 1.

CC PRINTS; PR01234; TNECROSISFCT.
CC ProDom; PD02012; TNF_subf; 1.
CC SMART; SM00207; TNF; 1.
CC PROSITE; PS00251; TNF_1; FALSE_NEG.
CC PROSITE; PS50049; TNF_2; 1.

CC Cytokine; Transmembrane; Glycoprotein; Signal-anchor;
CC Alternative splicing.
CC CHAIN 1 240

FT CHAIN 793 240 TUMOR NECROSIS FACTOR LIGAND SUPERF
FT CHAIN 793 240 TUMOR NECROSIS FACTOR LIGAND SUPERF
FT DOMAIN 1 37 MEMBER 14, SOLUBLE FORM.
FT TRANSMEM 38 58 CYTOPLASMIC (POTENTIAL).

FT DOMAIN 59 240 SIGNAL-ANCHOR (TYPE-II MEMBRANE PRO
FT SITE 82 83 (POTENTIAL).
FT SITE 82 83 EXTRACELLULAR (POTENTIAL).
FT DISULFID 154 187 CLEAVAGE (POTENTIAL).
FT CARBOHYD 102 102 POTENTIAL.

FT VARSPLIC 38 73 N-LINKED (GLCNAC. . .).
FT CONFLICT 120 120 Missing (in isoform 2).
FT CONFLICT 214 214 /FTid=VSP_006452.
FT CONFLICT 214 214 L -> V (IN REF. 4).
FT SEQUENCE 240 AA; 26351 MW; 49D0BF67E1390B39 CRC64;

Query Match 6.4%; Score 92; DB 1; Length 240;
Best Local Similarity 23.7%; Pred. No. 2.8;
Matches 44; Conservative 19; Mismatches 57; Indels 66; (

QY 29 QAQPAPMAARSQRRCRGPCTALLVPLALGLGLALACGLLLAVVLSGRAS
DB 16 QTDIPFTLGRSHRRSCSVARVGLGLLL-LIMGAGLAVQGVFLQLHWRIG----

QY 89 EPAQELVAEEDQDPSELNPOTESQDPAPFLNRLVLRPRRSPAPGRKTRARRAI
DB 67 -----EMV-----TRLPDGPAGSWEQLIQERS-----

QY 149 HPRPQDGAQGVDTGTVSGWEARINSSPLRYNRQI-----GEFIVT
DB 93 NPAHLTGANSLSLTG-----SCGPLLWETQLGLAFRLGLSYHDGALVVTI

QY 197 YLYQCV 202

1 V 147

STANDARD; PRT; 280 AA.

el. 41, Created

el. 41, Last sequence update)

el. 41, Last annotation update)

factor ligand superfamily member 6 (FAS antigen ligand)

OR CD95L.

quatus atys (Red-crowned mangabey) (Sooty mangabey).

azoa; Chordata; Craniata; Vertebrata; Euteleostomi;

eria; Primates; Catarrhini; Cercopitheidae;

1; Cercocebus.

N.A.

18; PubMed:11491535;

Sostik P., Mayne A.E., King C.L., Genain C.P.,

sari A.A.;

encing, and homology analysis of nonhuman primate

and co-stimulatory molecules.";

53:315-328(2001)

Cytokine that binds to TNFRSF6/FAS, a receptor that

the apoptotic signal into cells. May be involved in

T cell mediated apoptosis and in T cell development.

S-mediated apoptosis may have a role in the induction of

tolerance, in the antigen-stimulated suicide of mature

r both. Binding to the decoy receptor TNFRSF6B/DCR3

its effects (By similarity).

omotrimer (Probable).

R LOCATION: Type II membrane protein and secreted (By

oluble form derives from the membrane form by

c processing (By similarity).

: Belongs to the tumor necrosis factor family.

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il to license@isb-sib.ch).

AAK37606.1; -

4TSV.

08064; Fas_ligand.

06053; TNF abc.

06052; TNF family.

08983; TNF like.

03636; TNF_subf.

TNF; 1

1; FASLIGAND.

4; TNCRSISFCT.

12; TNF_subf; 1.

1; TNF; 1.

51; TNF 1; 1.

049; TNF2; 1.

osis; Transmembrane; Glycoprotein; Signal-anchor.

1 280

TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY

MEMBER 6, MEMBRANE FORM.

19 280

TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY

MEMBER 6, SOLUBLE FORM (BY SIMILARITY).

1 80

CYTOPLASMIC (POTENTIAL).

11 101

SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)

(POTENTIAL).

12 280

EXTRACELLULAR (POTENTIAL).

FT DOMAIN 4 69 PRO-RICH.

FT DOMAIN 45 64 POLY-PRO.

FT SITE 128 129 CLEAVAGE (BY SIMILARITY).

FT DISULFID 201 232 POTENTIAL.

FT CARBOHYD 183 183 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 249 249 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 259 259 N-LINKED (GLCNAC. . .) (POTENTIAL).

SQ SEQUENCE 280 AA; 31407 MW; 729EA60067B7D398 CRC64;

Query Match 6.4%; Score 92; DB 1; Length 280;

Best Local Similarity 20.8%; Pred. No. 3.4; Indels 74; Ga

Matches 62; Conservative 44; Mismatches 118;

QY 13 LPPLSLGSDGAVRQAPAPMAARRSQRRRGRGPGTALLVPLAL-----

Db 30 LPCPTSVRRRCQRRPPPPPPPLP-----PPPPPLPLPLPKKGN

QY 62 GLGLALACIGLLLVVSLGSRASLSAQPAQELVAEBDDQDPSSELNPQTESQDPAE

Db 79 GLCLDMFMFVLVALVGLG--LGMFQLFLQKEL-----AELRESTSQKTAS

QY 122 RLVRPRRSPKGRKTRARRAIAAHYEVHPRQDGAQAGVDGTVSGWEEA-RINSSE

Db 129 QIGHP---SPPEKKEQRK--VAHLTGKPNRSKMPLE-----WEDTYGIVLLK

QY 181 YNRQIGEFIVTRAGLYLYCQVHFDEGKA-----VYLKLD-----LLVDGVI

Db 175 YKK--GGLVINETGLYFYYSKYVF-RGQSCNPLSLSHKVMYNSKYPQDLVMEGKO

QY 226 CLUEPSATAASSLGQLRLCQVSGLLALRPGSLRLRTLPWAHLKAAPLTYFGLP{

Db 232 CWTGQWHAHSSYLGAFFNLSTDHLY-----VNVSELSLVNFESQ--TFGLYI

RESULT 14

NFC4 HUMAN

ID NFC4 HUMAN STANDARD; PRT; 902 AA.

AC Q14934;

DT 16-OCT-2001 (Rel. 40, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DE 15-MAR-2004 (Rel. 43, Last annotation update)

DE Nuclear factor of activated T-cells, cytoplasmic 4 (T cell

DE transcription factor NFAT3) (NF-ATc4) (NF-AT3).

GN NFATC4 OR NFAT3.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RC TISSUE=T-cell;

RX MEDLINE=95269130; PubMed=7749981;

RA Hoey T., Sun Y.-L., Williamson K., Xu X.;

RT "Isolation of two new members of the NF-AT gene family and function

RT characterization of the NF-AT proteins.";

RL Immunity 2:461-472(1995).

RN [2]

RP REVIEW.

RX MEDLINE=99189746; PubMed=10089876;

RA Crabtree G.R.;

RT "Generic signals and specific outcomes: signaling through Ca2+,

RT calcineurin, and NF-AT.";

RL Cell 96:611-614(1999).

CC -!- FUNCTION: Plays a role in the inducible expression of cytokine

CC genes in T cells, especially in the induction of the IL-2 and

CC 4 (By similarity).

CC -!- SUBUNIT: Member of the multicomponent NFATC transcription com

CC that consists of at least two components, a pre-existing

CC cytoplasmic component NFATC2 and an inducible nuclear component

CC NFATC1. Other members such as NFATC4, NFATC3 or members of the

CC activating protein-1 family, MAP, GATA4 and Cbp/p300 can also

CC the complex. NFATC proteins bind to DNA as monomers.

CC -!- SUBCELLULAR LOCATION: Cytoplasmic for the phosphorylated form

After activation that is controlled by calcineurin-phosphorylation. Rapid nuclear exit of NFATC is thought mechanism by which cells distinguish between sustained and transient calcium signals. The subcellular localization of NFATC is a key role in the gene transcription. NFATC1 is highly expressed in placenta, lung, kidney, and thymus. Not in peripheral blood lymphocytes. NFATC1 allows DNA-binding and NFATC1 interacts with API factors (By similarity). NFATC1 is phosphorylated by NFATC1-kinase; dephosphorylated by NFATC1 phosphatase (By similarity). NFATC1 belongs to the Rel/Dorsal family.

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AA79175.1; -
462; -
778; NFATC4.

3; P:transcription co-activator activity; TAS.

4; P:inflammatory response; TAS.

6; P:transcription from Pol II promoter; TAS.

007110; IG-like.

002909; IPT TIG.

00451; NF Rel dor.

008366; NFAT.

008367; P53-like.

; RHD; 1.

; TIG; 1.

89; NUCFACTORATC.

9; IPT; 1.

204; REL 1; FALSE NEG.

254; REL 2; 1.

regulation; Activator; Nuclear protein; DNA-binding; phosphorylation.

62 69 POLY-PRO.

14 119 CALCINEURIN-BINDING.

13 293 2 APPROXIMATE SP REPEATS.

13 229 SP 1.

77 293 SP 2 (APPROXIMATE).

97 304 POLY-PRO.

68 270 NUCLEAR LOCALIZATION SIGNAL.

30 437 DNA-BINDING.

72 674 NUCLEAR LOCALIZATION SIGNAL.

2 AA; 95472 MW; E59F15F7647A47C6 CRC64;

arity 6.4%; Score 92; DB 1; Length 902;

conservative 15; Mismatches 73; Indels 56; Gaps 8;

SLGSDGAVZQAQPPAPMAARRS-----QRRGRGREGTALLVPLALGLALAC 69

RGEDSLLLSAGPTASPRPASPCGRKRYSSSGTPSSA-----SPALSR 286

LAVSLGSRASLSAQEPQAEELVAEEDQPSSELPQTEESQDPAPFFLNRLVPRRS 129

-----SLGEGS-----EPPPPPL-PLARDPGSPGFEDVVGAPPAES 325

GRKTRARAI-----AHVEVHPRQDGAQAGVD-----GTVSG 167

TRTSSEQAVLPRSEEPASCNGLPLGABESVAPPGSRKEVAGMDYLAVPSPLA 385

RLNSSPL 179

RIGHHSPI 397

RESULT 15

MLH1_MOUSE STANDARD; PRT; 760 AA.
ID MLH1_MOUSE
AC Q9UK91; Q62454;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE DNA mismatch repair protein Mlh1 (MutL protein homolog 1).
GN MLH1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus;
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Testis;
RA Kumaran M., Rao M.R.S.;
RT "Cloning of the cDNA of the MutL homolog, MLH1 from mouse testis.
Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
[2]
RP SEQUENCE OF 1-151 FROM N.A.
RX MEDLINE=96270514; PubMed=8674118;
RA Edelmann W., Cohen P.E., Kane M., Lau K., Morrow B., Bennett S.,
RA Umar A., Kunkel T., Cattoretti G., Chaganti R., Pollard J.W.,
RA Kolodner R.D., Kucherlapati R.;
RT "Meiotic pachytene arrest in MLH1-deficient mice.";
RL Cell 85:1125-1134(1996).
CC -!- FUNCTION: Probably involved in the repair of mismatches in DN
CC -!- SUBUNIT: HETERODIMER OF MLH1 AND PMS2 OR MLH1 AND MLH3 (BY
CC SIMILARITY).
CC -!- SIMILARITY: Belongs to the DNA mismatch repair mutL/hexB fami
CC This SWISS-PROT entry is copyright. It is produced through a coll
CC between the Swiss Institute of Bioinformatics and the EMBL out
CC the European Bioinformatics Institute. There are no restriction
CC use by non-profit institutions as long as its content is in
CC modified and this statement is not removed. Usage by and for c
CC entities requires a license agreement (See <http://www.isb-sib.ch/>
CC or send an email to license@isb-sib.ch).

EMBL; AF250844; AAC52672.1; -
EMBL; U60872; AAC52672.1; -
EMBL; U59881; AAC52672.1; JOINED.
EMBL; U59882; AAC52672.1; JOINED.
EMBL; U59883; AAC52672.1; JOINED.
EMBL; U59884; AAC52672.1; JOINED.
HSP; P23367; 1BKN.
MGD; MGI:101938; Mlh1.
GO; GO:0000793; C:condensed chromosome; IDA.
GO; GO:0007060; P:meiotic chromosome segregation; IMP.
GO; GO:0007126; P:meiosis; IDA.
GO; GO:0007131; P:meiotic recombination; IMP.
GO; GO:0006298; P:mismatch repair; IDA.
InterPro; IPR002099; DNA_mis_repair.
Pfam; PF01119; DNA_mis_repair; 1.
Pfam; PF02518; HATPase_C; 1.
SMART; SM00387; HATPase_C; 1.
TIGRfams; TIGR00585; mutL; 1.
PROSITE; PS00058; DNA_MISMATCH_REPAIR_1; 1.
KW DNA repair.
SQ SEQUENCE 760 AA; 84679 MW; 173C809372A29186 CRC64;

Query Match 6.3%; Score 90.5; DB 1; Length 760;
Best Local Similarity 23.7%; Pred. No. 14;
Matches 70; Conservative 31; Mismatches 126; Indels 81; G

QY 13 LPLPRSLGSRDGGAVRQ-----AQPAPMAARRSQRGRGREGTALLVPLA
DB 436 LPAPAAEAASENLERESLMETSDAAQKAAPTSPGSRKRRH--EDSDVENVZ
QY 65 LALAC-----IGLLAVVSLGSRASLSAQEPAQEELVAEEDQDPSELNPO--TEES

06:25:24 2004

us-09-245-198a-4.rsp

1

YPRRIINLTSVLSEISERCHETLRE--ILRNHSFVGCVPQWALAQHOTKL 551
ILVPRRSAPKGRKTARRAIAHYEV-----HPRPGDGAQAGVDG 163
-----TTKJSEELFYQILIYDFANFGVLRRLSEPAFLDLAMLALDS 596
VEEARINSSPLRYNQIGEFIVTRAGLYLYGCVHFDEGKAVYKLIDLIVDG-- 220
VTEDDGPKGLABY---IVEFLKKAEMLADYESVEIDE--EGNLIIGLPLILDSYVP 652
--VIALR-----CLEEFSATAAS--SLGPQ--LRLCQVSGLLALRPSS 258
JPIFILRLATEVNVNDEEKECFESLSKECAMFYSIKQYILEESTLSGQQSDMEGST 712
CLEW 266
--FW 716

April 7, 2004, 17:45:19
secs

6:25:22 2004

us-09-245-198a-4.rag

GenCore version 5.1.6
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search, using sw model

il 7, 2004, 17:37:32 ; Search time 56.9116 Seconds
(without alignments)
1409.967 Million cell updates/sec

09-245-198A-4
4
SLLDFFESARRLPRLPSLG.....PWAHLKAAPETYGLFQVH 284

SUM62

op 10.0 , Gapext 0.5

6107 seqs, 282547505 residues

s satisfying chosen parameters: 1586107

th: 0

th: 2000000000

nimum Match 0%

ximum Match 100%

sting first 45 summaries

Geneseq 29Jan04: *

geneseq1980s: *

geneseq1990s: *

geneseq2000s: *

geneseq2001s: *

geneseq2002s: *

geneseq2003as: *

geneseq2003bs: *

geneseq2004s: *

the number of results predicted by chance to have a
than or equal to the score of the result being printed,
d by analysis of the total score distribution.

SUMMARIES

ry	ch	Length	DB	ID	Description
.0	284	2	Aaw47525	Homo sapi	
.8	249	2	Aay09369	Human tum	
.8	249	3	Aay95338	Human PRO	
.8	249	3	Aab07526	Amino aci	
.8	249	5	Aau86129	Human PRO	
.8	249	6	Abra42315	Human TWB	
.8	249	7	Adc35206	Human TNF	
.6	249	2	Aaw29745	TNF relat	
.6	249	4	Aae00891	Human TRE	
.6	249	7	Adc97712	Marine FL	
.8	273	4	Aau03499	TWEAK ext	
.5	208	2	Aaw93590	Human TNR	
.6	225	2	Aaw47524	Mus muscu	
.6	225	3	Aab07527	Amino aci	
.0	211	2	Aaw93591	Mouse TNR	
.8	189	2	Aaw29746	TNF relat	
.8	189	4	Aae00892	Human U14	
.7	146	4	Aae00895	Human TRE	
.0	325	4	Abb67553	Drosophil	
.0	409	5	Aau77718	Drosophil	
.5	211	3	Aay58216	Canine na	
.5	260	3	Aay58215	Canine CD	
.4	254	4	Aar64190	Human 4-1	
.4	254	2	Aaw26657	Human 4-1	
.4	254	5	Abb75953	Human cyt	

26	106.5	7.4	254	6	ABR42312	Hum
27	106.5	7.4	254	7	ADC35200	Hum
28	106.5	7.4	254	7	ADD18780	Hum
29	106	7.3	1428	3	AAy97033	Cas
30	105.5	7.3	406	5	AAU77717	Dro
31	104.5	7.2	779	5	ABB07845	Hum
32	104	7.2	409	5	AAU77716	Dro
33	104	7.2	1323	2	AAR55248	N-m
34	102	7.1	256	4	AAM25657	Hum
35	100.5	7.0	647	2	AAW04327	Rat
36	100	6.9	220	4	AA62340	Gp1
37	99	6.9	574	3	AAy97032	Cas
38	99	6.9	855	7	ADB96563	Hum
39	99	6.9	915	6	ABP56824	Hum
40	99	6.9	1008	4	AAW78891	Hum
41	99	6.9	1020	4	AAW79875	Hum
42	97.5	6.8	1560	7	ADE71264	Nov
43	96.5	6.7	1097	4	ABG25655	Nov
44	96.5	6.7	1631	4	ABG22481	Nov
45	96.5	6.7	19938	6	ABP76681	Str

ALIGNMENTS

RESULT 1
AAW47525
ID AAW47525 standard; protein; 284 AA.
XX
AC AAW47525;
XX
DT 21-JUL-1998 (first entry)
XX
DE Homo sapiens tumour necrosis factor related ligand (TRELL).
XX
KW TRELL; tumour necrosis factor related ligand; tnfr; treatment; can
autoimmune disease; immune system; stimulation; suppression;
KW graft rejection.
XX
OS Homo sapiens.
XX
PN WO9805783-Al.
XX
PD 12-FEB-1998.
XX
PF 07-AUG-1997; 97WO-US013945.
XX
PR 07-AUG-1996; 96US-0023541P.
PR 18-OCT-1996; 96US-0028515P.
PR 18-MAR-1997; 97US-0040820P.
XX
PA (BIOJ) BIOGEN INC.
PA (UYGE-) UNIV GENEVA FACULTY MEDICINE.
XX
PI Chicheportiche Y, Browning JL;
DR WPI: 1998-145619/13.
DR N-PSDB; AAV18600.
XX
PT Tumour necrosis factor related ligand - useful for, e.g. treating
auto-immune disease and immune responses to tissue grafts.
XX
PS Claim 12; Page 50-51; 69pp; English.
XX
CC The sequence is that of human tumour necrosis factor related ligar
(TRELL). TRELL or active fragments can be included with a carrier
pharmaceutical compositions to treat cancer, autoimmune diseases c
immune responses to tissue grafts, or to stimulate or suppress the
system. It is useful to screen for TRELL receptors, by labelling a
detectable label and screening compositions for binding. Agents
interfering with TRELL-receptor binding can also be screened for,
then be administered, optionally with interferon- gamma, to induce
death or treat, suppress or alter immune responses (especially inv

ccinoma cells) involving a signal pathway between TRELL and its coding sequence can be used in gene therapy for TRELL-defects in mammals (especially humans), e.g. tumours, inflammatory diseases or inherited genetic disorders, by to cells, and expressing, therapeutically effective amounts e.g. a virus comprising a gene encoding TRELL. It may also the preparation of prepare probes for screening etic DNAs for TRELL-encoding sequences and for antisense

AA;
100.0%; Score 1444; DB 2; Length 284;
arity 100.0%; Pred. No. 2.4e-129;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
DFEISARRLPLPSLGRDGGAVTQAPPPAPMAARRSQRGRGEGPTALLVPLA 60
DFEISARRLPLPSLGRDGGAVTQAPPPAPMAARRSQRGRGEGPTALLVPLA 60
LALACLGLLLAVSLGSRASLSAQEELVAEEDQDPSELNPQTESQDPAPFL 120
LALACLGLLLAVSLGSRASLSAQEELVAEEDQDPSELNPQTESQDPAPFL 120
RPRSAPKGRKTRARRATAAHYEVHPRPGDGAQAGVDGTGVSWEARINSSPLR 180
RPRSAPKGRKTRARRATAAHYEVHPRPGDGAQAGVDGTGVSWEARINSSPLR 180
IGFEIVTRAGLYLYCQVHFDEGKAVYKLLDLDVGLALRCLEEFSAATASSLGP 240
IGFEIVTRAGLYLYCQVHFDEGKAVYKLLDLDVGLALRCLEEFSAATASSLGP 240
CQVSGLLALRPGSSLRINTLPWAHLKAAPFLTYFGLFQVH 284
CQVSGLLALRPGSSLRINTLPWAHLKAAPFLTYFGLFQVH 284

dard; protein; 249 AA.

(first entry)

necrosis factor Apo-3 ligand protein sequence.

necrosis factor; Apo-3 ligand; lymphotoxin; apoptosis;
ndent transcription; JNK/SAPK-dependent responses; cancer.

98WC-US021407.

97US-0062037P.

97US-0069862P.

TECH INC.

Marsters SA, Pitti R;

1982/24.

5000.

3- ligand (a tumor necrosis factor) homologue.

1; 74pp; English.

sequence represents a human tumour necrosis factor (TNF) and

CC lymphotoxin homologue designated Apo-3 ligand. Apo-3 ligand has
CC cytostatic activity. Apo-3 ligand can be used to induce apoptosis
CC mammalian cancer cells, to induce NF-kappaB-dependent transcripti
CC to induce JNK/SAPK-dependent responses in mammalian cells
XX
SQ Sequence 249 AA;

Query Match 87.8%; Score 1268; DB 2; Length 249;
Best Local Similarity 100.0%; Pred. NO. 1.3e-112;
Matches 249; Conservative 0; Mismatches 0; Indels 0; G

Qy 36 MAARRSORRRGRGEGPTALLVPLALGLGLALACLGALLAVSLGSRASLSAQEPA
Db 1 MAARRSORRRGRGEGPTALLVPLALGLGLALACLGALLAVSLGSRASLSAQEPA
Qy 96 VAEDQDPSELNPQTESQDPAPFLNLRVPRRSAPKGRKTRARRATAAHYEVHPR
Db 61 VAEDQDPSELNPQTESQDPAPFLNLRVPRRSAPKGRKTRARRATAAHYEVHPR
Qy 156 GAQAGVDGTGVSWEARINSSPLRYNRQIGFEIVTRAGLYLYCQVHFDEGKAVY
Db 121 GAQAGVDGTGVSWEARINSSPLRYNRQIGFEIVTRAGLYLYCQVHFDEGKAVY
Qy 216 LLVDGVLALRCLEEFSAATASSLGPQLRCCQVSGLLALRPGSSLRINTLPWAHLKA
Db 181 LLVDGVLALRCLEEFSAATASSLGPQLRCCQVSGLLALRPGSSLRINTLPWAHLKA
Qy 276 TYFGLFQVH 284
Db 241 TYFGLFQVH 249

RESULT 3

AAAY95338

ID AAAY95338 standard; protein; 249 AA.

AC AAAY95338;

XX 25-SEP-2000 (first entry)

XX Human PRO207 antitumour protein.

XX PRO207; human; antitumour; tumour; therapy; cytostatic; breast ca
KW ovarian cancer; renal cancer; colorectal cancer; uterine cancer;
KW prostate cancer; lung cancer; bladder cancer;
KW central nervous system cancer; melanoma; leukaemia; neoplasia.

OS Homo sapiens.

XX Key Location/Qualifiers

FH Peptide 1..40

FT /label= Signal_peptide

FT Modified-site 10..14

FT /note= "amidation"

FT Peptide 24..35

FT /note= "prokaryotic membrane lipoprotein lipid"

FT Modified-site 27..33

FT /note= "N-myristoylation"

FT Modified-site 29..35

FT /note= "N-myristoylation"

FT Modified-site 36..42

FT /note= "N-myristoylation"

FT Protein 41..249

FT /label= PRO207

FT Modified-site 45..51

FT /note= "N-myristoylation"

FT Modified-site 97..101

FT /note= "amidation"

FT Modified-site 118..124

FT /note= "N-myristoylation"

FT Modified-site 121..127

FT /note= "N-myristoylation"

FT Modified-site 125..131

/note= "N-myristoylation"
128. .134
/note= "N-myristoylation"
139. .143
/note= "Asn is N-glycosylated"

99WO-US028565.

98US-0113296P.

99WO-US005028.

98US-0130232P.

98US-0131443P.

98US-0134287P.

98US-0144758P.

98US-0145698P.

99WO-US021090.

99WO-US021547.

ECH INC.

Goddard A., Godowski PJ, Gurney AL, Marsters SA;

tti RM, Wood WI;

58/38.

17.

ion to inhibit neoplastic cell growth or for treating tumor
ries polypeptides PRO179, PRO207, PRO320, PRO219, PRO221,
, PRO301, PRO526, PRO362, PRO356, PRO509 or PRO866.

4; 172pp; English.

quence is that of human antitumor protein PRO207, as
foetal kidney cDNA clone (see AAA49717). PRO207 shows
ence identity to tumour necrosis factor family members,
an lymphotoxin-beta (23.4%) and human CD40 ligand (19.8%).
216. A claimed method for inhibiting the growth of a tumour
exposing the tumor cell to PRO179, PRO207, PRO320, PRO219,
, PRO328, PRO301, PRO526, PRO362, PRO356, PRO509 or PRO866
49), their agonists or chimeric polypeptides incorporating
ur is especially a cancer selected from breast, ovarian,
tal, uterine, prostate, lung, bladder and central nervous
melanoma and leukaemia. Methods for the recombinant
the antitumor proteins are also provided

A;

87.8%; Score 1268; DB 3; Length 249;
rity 100.0%; Pred. No. 1.3e-112; Indels 0; Gaps 0;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
SQRRRGRGEGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPAQEL 95
SQRRRGRGEGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPAQEL 60
QDPSELNPQTEESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPRPGQD 155
QDPSELNPQTEESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPRPGQD 120
VDGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCOVHDEGKAVYKLD 215
VDGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCOVHDEGKAVYKLD 180
VLAALRCLEEFSAASSLGPQLRCQVSGLLALPGSSLRIRTLTPWAHLKAPFL 275
VLAALRCLEEFSAASSLGPQLRCQVSGLLALPGSSLRIRTLTPWAHLKAPFL 240
FQVH 284
|||

Db 241 TYFGLFQVH 249

RESULT 4
AAB07526
ID AAB07526 standard; protein; 249 AA.
XX AAB07526;
AC AAB07526;
XX 20-OCT-2000 (first entry)
XX 20-OCT-2000 (first entry)
XX Amino acid sequence of a soluble recombinant human TWEAK protein.
DE TWEAK protein; immunological disorder; immune response; inflammati
KW TWEAK blocking agent; autoimmune disease; organ transplant rejecti
KW Graft-versus-Host disease; GVHD; lymphoid cell malignancy; shock;
XX Homo sapiens.
OS WO200042073-A1.
XX 20-JUL-2000.
PD 14-JAN-2000; 2000WO-US001044.
XX 15-JAN-1999; 99US-0116168P.
PR (BIOJ) BIOGEN INC.
PA Renmert P;
XX WPI; 2000-476036/41.
DR Preventing and treating immune responses using modulators, especia
PT antibodies, of TWEAK, TWEAK receptors and TWEAK ligands, useful fo
PT treating e.g. inflammation and graft versus host disease.
XX Disclosure; Fig 1; 45pp; English.
XX The present sequence represents a TWEAK protein. The specification
describes a method for preventing or treating an immunological dis
and/or inhibiting an immune response in an animal. The method comp
administering a TWEAK blocking agent. The method may be used for
preventing and treating immune disorders associated with inappropi
expression and/or activity of TWEAK. These disorders include autoi
diseases, acute and chronic inflammation, organ transplant rejecti
Graft-versus-Host disease (GVHD), lymphoid cell malignancies, sept
other forms of shock, loss of immune responsiveness (as seen in hu
immunodeficiency virus (HIV) infections) and failure of the immune
response to tumour growth
XX Sequence 249 AA;
Query Match 87.8%; Score 1268; DB 3; Length 249;
Best Local Similarity 100.0%; Pred. No. 1.3e-112; Indels 0; G
Matches 249; Conservative 0; Mismatches 0; Indels 0; G
QY 36 MAARSQRRGRGEGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPAQ
Db 1 MAARSQRRGRGEGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPAQ
QY 96 VAEEDQDPSELNPQTEESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPR
Db 61 VAEEDQDPSELNPQTEESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPR
QY 156 GAQAGVDGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCOVHDEGKAVY
Db 121 GAQAGVDGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCOVHDEGKAVY
QY 216 LIAVDGVLALRCLEEFSAASSLGPQLRCQVSGLLALPGSSLRIRTLTPWAHLKAF
Db 181 LIAVDGVLALRCLEEFSAASSLGPQLRCQVSGLLALPGSSLRIRTLTPWAHLKAF

1.FQVH 284
1.FQVH 249

ard; protein; 249 AA.

(first entry)
polypeptide.

enign tumour; malignant tumour; lymphoid malignancy;
uronal disorder; stromal disorder; blastocoeleic disorder;
disorder; immune disorder; angiogenic disorder; cytostatic;
ve.

1.

2000WO-US003565.

99WO-US005028.
99US-0123972P.
99US-0133459P.
99WO-US012252.
99US-0140650P.
99US-0140653P.
99US-0144758P.
99US-0145698P.
99US-0146222P.
99US-0149395P.
99US-0151689P.
99WO-US020111.
99WO-US021090.
99WO-US028313.
99WO-US028301.
99WO-US028634.
2000WO-US000219.

TECH INC.

Goddard A, Godowski PJ, Gurney AL, Hillan KJ;
Pan J, Pitti RM, Roy MA, Smith V, Stone DM;
Wood WI;

567/26.
255.

nucleic acids encoding PRO polypeptides, useful for treating
ignant tumors, leukemias and lymphoid malignancies,
angiogenic and immunologic disorders.

4; 302pp; English.

vention relates to the isolation of novel human PRO
and the polynucleotide sequences encoding them. The PRO
agonists, antagonists or anti-PRO antibodies are useful for
gn or malignant tumours (e.g. renal, kidney, bladder, such
leukaemias and lymphoid malignancies, other disorders such
glial, astrocytal, hypothalamic, glandular, macrophagal,
blastocoeleic disorders, inflammatory, immune and angiogenic
e polynucleotide sequences are also useful in gene therapy.
16162 represent the human PRO polypeptides of the invention

AA;

Query Match 87.8%; Score 1268; DB 5; Length 249;
Best Local Similarity 100.0%; Pred. No. 1.3e-112;
Matches 249; Conservative 0; Mismatches 0; Indels 0; G

QY 36 MAARRSQRGRGEGPTALLVPLALGLGLALACLGILLAVVSLGSRASLSAQEPA
Db 1 MAARRSQRGRGEGPTALLVPLALGLGLALACLGILLAVVSLGSRASLSAQEPA
QY 96 VAEEDQDPSELNPQTEESQDPAPFLNRLVRPRSPKGRKTRARRAIAAHYEVHPR
Db 61 VAEEDQDPSELNPQTEESQDPAPFLNRLVRPRSPKGRKTRARRAIAAHYEVHPR
QY 156 GAQAGVDGTVSGWEHARINSSPLRYNRQIGEFIVTRAGLYLYLYCQVHFDEGKAVY
Db 121 GAQAGVDGTVSGWEHARINSSPLRYNRQIGEFIVTRAGLYLYLYCQVHFDEGKAVY
QY 216 LLDVGVIALRCLFEFSATAASSLGPQLRCLQVSGLLALRPGSSLRIRTLFWAHLKA
Db 181 LLDVGVIALRCLFEFSATAASSLGPQLRCLQVSGLLALRPGSSLRIRTLFWAHLKA
QY 276 TYFGLFQVH 284
Db 241 TYFGLFQVH 249

RESULT 6

ABR42315
ID ABR42315 standard; protein; 249 AA.

AC ABR42315;

DT 11-AUG-2003 (first entry)

DE Human TWEAK protein.

KW Human; TWEAK; tumour necrosis factor; ligand; cytostatic;
immunomodulator; osteopathic.

OS Homo sapiens.

PN WO2003040307-A2.

PD 15-MAY-2003.

PF 25-JUL-2002; 2002WO-US023782.

PR 27-JUL-2001; 2001US-0307838P.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Hilbert DH, Rosen CA;

DR WPI; 2003-430659/40.

DR N-PSDB; ACC57901.

XX New heteromultimeric complex having a first polypeptide member of
tumor necrosis factor (TNF) ligand family, and a second different
of TNF ligand family, useful for treating cancer, osteoporosis or
autoimmune disease.

PS Disclosure; Page 368-369; 388pp; English.

XX The present sequence is the protein sequence for human TWEAK prot
invention relates to compositions comprising heterotrimeric compl
tumor necrosis factor (TNF) ligand family members, and their use
detection, prevention and treatment of disease. In one embodiment
heterotrimeric complex comprises full-length or extracellular por
TWEAK and full-length or extracellular portions of other TNF lig
family members, preferably VEGI or VEGI-SV. The heterotrimeric co
of the invention are useful for treating an autoimmune disease, c
osteoporosis, and particularly for inhibiting cancer cell prolif
increasing B cell proliferation, or inducing apoptosis of T cells

XX

87.8%; Score 1268; DB 6; Length 249;
 city 100.0%; Pred. No. 1.3e-112; Indels 0; Gaps 0;
 iservative 0; Mismatches 0; Indels 0; Gaps 0;
 SQRGRGPGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPQEL 95
 SQRGRGPGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPQEL 60
 QDSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPRQD 155
 QDSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPRQD 120
 VDGTVSGWEERINSSPLRYNROI GEFIVTRAGLYLYYLCQVHDEGKAVYLKLD 215
 VDGTVSGWEERINSSPLRYNROI GEFIVTRAGLYLYYLCQVHDEGKAVYLKLD 180
 VLALRCLEEPSATAASLGQRLCQVSGLLALRPGSSLRIRTL PWAHLKAAPPL 275
 VLALRCLEEPSATAASLGQRLCQVSGLLALRPGSSLRIRTL PWAHLKAAPPL 240
 FQVH 284
 FQVH 249

ard; protein; 249 AA.

first entry)

nd family member #12.

necrosis factor; TNF ligand; endokine alpha;
 resorption disorder; osteoporosis; Paget's disease;
 fication.

1.

002US-00218547.

001US-0312542P.

001US-0330761P.

C A.

LI B.

Rosen CA, Nardelli B;

172/66.

05.

alpha gene useful for preparing a composition for treating a
 iated with excessive or insufficient bone resorption e.g.,
 Paget's disease or arterial calcification.

EQ ID NO 24; 145pp; English.

relates to an isolated nucleic acid molecule encoding a
 is factor family ligand. A composition comprising the
 body or its fragment is used for treating an individual in
 ased level of endokine alpha activity. The endokine alpha
 resent in a heterotrimeric complex is used for treating an

CC individual having a disorder associated with excessive bone resorp
 e.g. osteoporosis, Paget's disease or arterial calcification. Trea
 CC individual having a disorder associated with insufficient bone res
 CC comprises administering an endokine alpha antagonist, which is the
 CC antibody that binds specifically to endokine alpha polypeptide. Th
 CC present sequence represents the amino acid sequence of a tumour ne
 CC factor family ligand.

Sequence 249 AA;

Query Match 87.8%; Score 1268; DB 7; Length 249;
 Best Local Similarity 100.0%; Pred. No. 1.3e-112;
 Matches 249; Conservative 0; Mismatches 0; Indels 0; Ga

QY 36 MAARRSQRRGRGPGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPAC
 Db 1 MAARRSQRRGRGPGTALLVPLALGLALACGLLLAVVSLGSRASLSAQEPAC
 QY 96 VAERDQDSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPRF
 Db 61 VAERDQDSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHYEVHPRF
 QY 156 GQAQGVDTGTVSGWEERINSSPLRYNROI GEFIVTRAGLYLYYLCQVHDEGKAVYI
 Db 121 GQAQGVDTGTVSGWEERINSSPLRYNROI GEFIVTRAGLYLYYLCQVHDEGKAVYI
 QY 216 LLVDGVLALRCLEEPSATAASLGQRLCQVSGLLALRPGSSLRIRTL PWAHLKAF
 Db 181 LLVDGVLALRCLEEPSATAASLGQRLCQVSGLLALRPGSSLRIRTL PWAHLKAF
 QY 276 TYFGLFQVH 284
 Db 241 TYFGLFQVH 249

RESULT 8

AAW29745

ID AAW29745 standard; protein; 249 AA.

XX AC AAW29745;

DT 27-OCT-1998 (first entry)

DE TNF related endothelium proliferative agent protein.

XX TNF; endothelium proliferative agent; TREPA; wound healing; cance.
 XX tissue grafting; vascularisation; apoptosis; autoimmune; birth co

OS HOMO sapiens.

XX WO9835061-A2.

XX 13-AUG-1998.

XX 12-FEB-1998; 98WO-US002859.

XX 12-FEB-1997; 97US-00798692.

XX 10-FEB-1998; 98US-00021706.

XX (ABBO) ABBOTT LAB.

XX Wiley SR;

XX WPI; 1998-447255/38.

XX N-PSDB; AAV47613.

XX Detecting nucleic acid encoding TREPA - useful for diagnosis and
 PT treatment of autoimmune disease, tumours and inflammation.

XX Claim 16; Page 123-4; 142pp; English.

XX The TNF-related endothelium proliferative agent (TREPA), or its
 CC activators or agonists, are used to treat a deficit of TREPA, e.g

healing or tissue grafting, by promoting vascularisation, a apoptosis for treating cancer and eliminating autoreactive n adjunct to cancer chemotherapy or antiviral treatment. s can also be used to target cytotoxic agents or for action of the corresponding receptor, the nucleic acid for used to transform tumour cells to render them more TREPA and to screen for TREPA mimics. Ribozymes, antisense ies or peptides, are used to treat TREPA-associated . tumours and metastases (by inhibiting vascularisation), or a wide range of autoimmune conditions, conditions ormal stimulation of epithelial cells (e.g. is), for birth control (inhibiting ovulation and placental other angiogenic conditions (e.g. ulcers)

AA;

87.6%; Score 1265; DB 2; Length 249;
arity 99.6%; Pred. No. 2.5e-112;
onservative 1; Mismatches 0; Indels 0; Gaps 0;

RSQRRRGEPGCTALLVPLALGLALACGLALLAVVSLGSRASLSAQEPQEL 95
RSQRRRGEPGCTALLVPLALGLALACGLALLAVVSLGSRASLSAQEPQEL 60
RSQRRRGEPGCTALLVPLALGLALACGLALLAVVSLGSRASLSAQEPQEL 120

DQDPSLNPTQEEQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHVEVHPRPQD 155
DQDPSLNPTQEEQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHVEVHPRPQD 120
DQDPSLNPTQEEQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHVEVHPRPQD 120

GVDTGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVYLKLD 215
GVDTGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVYLKLD 180
GVDTGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVYLKLD 180

GVALRCLEEFSAATAASLGQPLRCQVSGLLALRPGSSLRITLPAHLKAAPFL 275
GVALRCLEEFSAATAASLGQPLRCQVSGLLALRPGSSLRITLPAHLKAAPFL 240
GVALRCLEEFSAATAASLGQPLRCQVSGLLALRPGSSLRITLPAHLKAAPFL 240

LFQVH 284

LFQVH 249

andard; protein; 249 AA.

(first entry)

(TNF related endothelium proliferative agent).

: necrosis factor; TNF; angiogenesis; wound healing; TREPA;
ndothelium proliferative agent; tumour; metastasis;
nerary.

Location/Qualifiers

98..249

/label= Extracellular_domain

98US-00105343.

97US-00798692.

98US-00021706.

TT LAB.

XX WPI; 2001-280760/29.
DR N-PSDB; AAD04350.
XX Inducing angiogenesis in mammal at desired sites for promoting wo
PT healing, by administering soluble fragment of extracellular domai
PT tumor necrosis factor related endothelium proliferative agent pro
XX Claim 1; Col 75-76; 53pp; English.

XX The present invention relates to extracellular signal molecules,
CC particularly members of tumour necrosis factor (TNF) family molec
CC designated as TREPA (TNF related endothelium proliferative agent)
CC Soluble biologically active TREPA are used to treat TREPA-associ
CC diseases, tumours or metastases. TREPA is used for inducing angic
CC in human for promoting wound healing and for vascularising grafted
CC for successful grafting and to promote tissue grafts. The present
CC acid sequence is clone ID #690050 human TREPA

SQ Sequence 249 AA;

Query Match 87.6%; Score 1265; DB 4; Length 249;
Best Local Similarity 99.6%; Pred. No. 2.5e-112;
Matches 248; Conservative 1; Mismatches 0; Indels 0; C

QY 36 MAARRSORRRGEPGCTALLVPLALGLALACGLALLAVVSLGSRASLSAQEPF
DB 1 MAARRSQRRRGEPGCTALLVPLALGLALACGLALLAVVSLGSRASLSAQEPF

QY 96 VAEDQDPSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHVEVHPF
DB 61 VAEDQDPSELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAIAAHVEVHPF

QY 156 QAQAGVDGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVI
DB 121 QAQAGVDGTVSGWEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVI

QY 216 LLVDGVIALRCLEEFSAATAASLGQPLRCQVSGLLALRPGSSLRITLPAHLKI
DB 181 LLVDGVIALRCLEEFSAATAASLGQPLRCQVSGLLALRPGSSLRITLPAHLKI

QY 276 TYFGLFQVH 284

DB 241 TYFGLFQVH 249

RESULT 10

ADC97712

ID ADC97712 standard; protein; 249 AA.

XX ADC97712;

XX 15-JAN-2004 (first entry)

XX Murine FL-TWEAK.

XX Murine; FL-TWEAK; TNF relatedness and weak ability to induce cel.
KW TNF; Tumour Necrosis Factor; TWEAK; fibrosis; cardiac disease;
KW liver disease; lung disease; kidney disease; skin disease;
KW skeletal muscle disease; adipose tissue disease;
KW gastrointestinal tract disease; pancreatic disease;
KW reproductive organ disease; neural disease; cartilage disease;
KW bone disease; connective tissue disease; cellular death; hepatot;
KW dermatological; gastrointestinal; osteopathic.

XX Mus sp.

XX WO2003086311-A2.

XX 23-OCT-2003.

XX 09-APR-2003; 2003WO-US011350.

XX

hard; protein; 208 AA.

(first entry)

rotein.

is factor receptor; signal transducer molecule; TNF; APO4; abnormality; gestational abnormality; prostate cancer; PO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy; disease; main; immunogen; antibody preparation; breast carcinoma; man.

98WO-US018393.

97US-00924634.

WASHINGTON.

1191/17.
1424.

rosis Factor family receptor polypeptides and ligands - gnosis and treatment of prostate cancer and developmental abnormalities.

13A; 156pp; English.

on describes isolated Tumor Necrosis Factor (TNF) family peptides: APO4, APO6, APO8 and APO9 or their active id isolated TNF related ligands 1 and 3 (TNRL1 and TNRL3) or fragments. APO4 is useful for diagnosing prostate cancer by levels of APO4 in an individual. Prostate cancer can also be 3 APO4 selective binding agents linked to a therapeutic polypeptides are also useful for identifying selective ts, useful in diagnosis/treatment of disease by binding of a polypeptide/active fragment which is extracellular, or the cell surface. The binding is preferably performed in clypeptides/ active fragments are also useful for screening and antagonists by binding and observing the change in APO4 fective pharmacological agents useful in diagnosis or disease are also identified using APO4 polypeptides/active d APO4 signal transducer molecules that specifically interact laemic domain of APO4 and detecting a change in level of APO4 e method is performed in vivo or in vitro. APO polypeptides ul as immunogens for preparing antibodies. APO4 is also gnosis/treatment of developmental or gestational s. APO8 was transfected to human breast carcinoma cell line nduced apoptosis

AA;

73.5%; Score 1062; DB 2; Length 208;

larity 99.5%; Pred. No. 4.9e-93;

Conservative 0; Mismatches 1; Indels 0; Gaps 0;

GSASLSAQEPAQELVAEEQDDPSLNPTQESQDPAPFLNLRPRRSAPKGRKT 136

GSASLSAQEPAQELVAEEQDDPSLNPTQESQDPAPFLNLRPRRSAPKGRKT 60

RAIAAHYVHPRPGDGAQAGVDGTSGWEARINSSPLRYNQIGEFIVTRAGLY 196

RAIAAHYVHPRPGDGAQAGVDGTSGWEARINSSPLRYNQIGEFIVTRAGLY 120

Qy 197 YLYQVHFDEGKAVYKLDLLVDGVLAIRCLEEFSAATASSLGQQLRCQVSGLLA
Db 121 YLYQVHFDEGKAVYKLDLLVDGVLAIRCLEEFSAATASSLGQQLRCQVSGLLA

Qy 257 SSLRIRTLPAHLKAAPFLTYEGLFQVH 284

Db 181 SSLRIRTLPAHLKAAPFLTYEGLFQVH 208

RESULT 13

AAW47524

ID AAW47524 standard; protein; 225 AA.

XX AC AAW47524;

XX DT 21-JUL-1998 (first entry)

XX DE Mus musculus tumour necrosis factor related ligand (TRELL).

XX KW TRELL; tumour necrosis factor related ligand; tnfr; treatment; car

XX KW auto-immune disease; immune system; stimulation; suppression;

XX KW graft rejection.

XX OS Mus musculus.

XX FH Key

XX FT Domain

XX FT Location/Qualifiers

XX FT 1..21

XX FT /note= "hydrophobic, transmembrane domain"

XX PN WO9805783-A1.

XX PD 12-FEB-1998.

XX PF 07-AUG-1997; 97WO-US013945.

XX PR 07-AUG-1996; 96US-0023541P.

XX PR 18-OCT-1996; 96US-0028515P.

XX PR 18-MAR-1997; 97US-0040820P.

XX PA (BIOJ) BIOGEN INC.

XX PA (UYGE-) UNIV GENEVA FACULTY MEDICINE.

XX PI Chicheportiche Y, Browning JL;

XX DR WPI; 1998-145619/13.

XX DR N-PSDB; AAV18599.

XX PT Tumour necrosis factor related ligand - useful for, e.g. treatin

XX PT auto-immune disease and immune responses to tissue grafts.

XX PS Claim 12; Page 48-50; 69pp; English.

XX CC The sequence is that of mouse tumour necrosis factor related lig

XX CC (TRELL). TRELL or active fragments can be included with a carrie

XX CC pharmaceutical compositions to treat cancer, autoimmune diseases

XX CC immune responses to tissue grafts, or to stimulate or suppress

XX CC system. It is useful to screen for TRELL receptors, by labelling

XX CC detectable label and screening compositions for binding. Agents

XX CC interfering with TRELL-receptor binding can also be screened for

XX CC then be administered, optionally with interferon- gamma, to indu

XX CC death or treat, suppress or alter immune responses (especially i

XX CC human adenocarcinoma cells) involving a signal pathway between

XX CC its receptor. It's coding sequence can be used in gene therapy f

XX CC related disorders in mammals (especially humans), e.g. tumours,

XX CC autoimmune and inflammatory diseases or inherited genetic disord

XX CC introducing into cells, and expressing, therapeutically effectiv

XX CC of a vector, e.g. a virus comprising a gene encoding TRELL. It n

XX CC be of use in the preparation of prepare probes for screening

XX CC natural/synthetic DNAs for TRELL-encoding sequences and for anti

XX CC therapy

XX CC Sequence 225 AA;

XX SQ

70.6%; Score 1020; DB 2; Length 225;
 88.8%; Pred. No. 5.7e-89;
 9; Mismatches 16; Indels 0; Gaps 0;
 ALACGLLLAVVSLGSRASLSAQEPQAQELVAEEDQDPSELNPQTESQDP
 ALACGLLLVVSLSGWSATLSAQEPSELTAEDRRPELNPQTESQDVV
 PRSAPKGRKTRARRAIAAHYEVHPRPGDGAQAGVDTGVSWEARINSS
 PRSAPKGRKTRARRAIAAHYEVHPRPGDGAQAGVDTGVSWEETKINSS
 GEFFIVTRAGLYLYCQVHDEGKAVYLLKDLVGVLAALCLLEFSATAAS
 GEFTVIRAGLYLYCQVHDEGKAVYLLKDLVGVLAALCLLEFSATAAS
 QVSGLLALRPGSSLRIRTPWAHLKAAPFLTYFGLFQVH 284
 QVSGLLALRPGSSLRIRTPWAHLKAAPFLTYFGLFQVH 225

iard; protein; 225 AA.

(first entry)

quence of a soluble recombinant murine TWEAK protein.

; immunological disorder; immune response; inflammation;
 3 agent; autoimmune disease; organ transplant rejection;
 host disease; GVHD; lymphoid cell malignancy; shock; tumour.

1.

2000WO-US001044.

98US-0116168P.

N INC.

036/41.

d treating immune responses using modulators, especially
 f TWEAK, TWEAK receptors and TWEAK ligands, useful for
 inflammation and graft versus host disease.

ig 1; 45pp; English.

sequence represents a TWEAK protein. The specification
 method for preventing or treating an immunological disorder
 ting an immune response in an animal. The method comprises
 a TWEAK blocking agent. The method may be used for
 id treating immune disorders associated with inappropriate
 id/or activity of TWEAK. These disorders include autoimmune
 ire and chronic inflammation, organ transplant rejection,
 Host disease (GVHD), lymphoid cell malignancies, septic and
 f shock, loss of immune responsiveness (as seen in human
 ency virus (HIV) infections) and failure of the immune
 tumour growth

AA;

70.6%; Score 1020; DB 3; Length 225;

Best Local Similarity 88.8%; Pred. No. 5.7e-89;
 Matches 199; Conservative 9; Mismatches 16; Indels 0; G
 QY 61 LGIGLALACGLLLAVVSLGSRASLSAQEPQAQELVAEEDQDPSELNPQTESQDP
 Db 2 LSJGLALACGLLLAVVSLGWSATLSAQEPSELTAEDRRPELNPQTESQDV
 QY 121 NLRVPRRSPKGRKTRARRAIAAHYEVHPRPGDGAQAGVDTGVSWEARINSS
 Db 62 EQLVRRSPKGRKTRARRAIAAHYEVHPRPGDGAQAGVDTGVSWEETKINSS
 QY 181 YNQIGEFFIVTRAGLYLYCQVHDEGKAVYLLKDLVGVLAALCLLEFSATAAS
 Db 122 YDRQIGEFFIVTRAGLYLYCQVHDEGKAVYLLKDLVGVLAALCLLEFSATAAS
 QY 241 QRLCQVSGLLALRPGSSLRIRTPWAHLKAAPFLTYFGLFQVH 284
 Db 182 QRLCQVSGLLALRPGSSLRIRTPWAHLKAAPFLTYFGLFQVH 225

RESULT 15

AAW93591
 ID AAW93591 standard; protein; 211 AA.

XX AC AAW93591;

XX XX 18-JUN-1999 (first entry)

XX XX Mouse TNRL3 protein.

XX XX Tumour necrosis factor receptor; signal transducer molecule; TNF;
 KW developmental abnormality; gestational abnormality; prostate ca
 KW APO6; APO8; APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy;
 KW cytoplasmic domain; immunogen; antibody preparation; breast carci
 KW apoptosis; mouse.

XX OS Mus sp.

XX XX WO9911791-A2.

XX XX 11-MAR-1999.

XX PF 04-SEP-1998; 98WO-US018393.

XX PR 05-SEP-1997; 97US-00924634.

XX XX (UNIV) UNIV WASHINGTON.

XX XX Chaudhary PM;

XX DR WPI; 1999-205191/17.

XX DR N-PSDB; AAX23425.

XX XX New Tumor Necrosis Factor family receptor polypeptides and ligand
 PT useful for diagnosis and treatment of prostate cancer and develop
 PT or gestational abnormalities.

XX PS Claim 40; Fig 13B; 156pp; English.

XX CC This invention describes isolated Tumor Necrosis Factor (TNF) fan
 CC receptor polypeptides: APO4, APO6, APO8 and APO9 or their active
 CC fragments, and isolated TNF related ligands 1 and 3 (TNRL1 and TN
 CC their active fragments. APO4 is useful for diagnosing prostate ca
 CC determining levels of APO4 in an individual. Prostate cancer can
 CC treated using APO4 selective binding agents linked to a therapeutic
 CC moiety. APO4 polypeptides are also useful for identifying select
 CC binding agents, useful in diagnosis/treatment of disease by bindi
 CC agents to the polypeptide/active fragment which is extracellular,
 CC expressed on the cell surface. The binding is preferably perform
 CC vivo. APO4 polypeptides/ active fragments are also useful for sc
 CC for agonists and antagonists by binding and observing the change
 CC activity. Effective pharmacological agents useful in diagnosis o
 CC treatment of disease are also identified using APO4 polypeptides,

AP04 signal transducer molecules that specifically interact
asemic domain of APO4 and detecting a change in level of APO4
method is performed in vivo or in vitro. APO polypeptides
l as immunogens for preparing antibodies. APO4 is also
agnosis/treatment of developmental or gestational
. APO8 was transfected to human breast carcinoma cell line
duced apoptosis

AA;

67.0%; Score 968; DB 2; Length 211;
arity 89.1%; Pred. No. 4.8e-84;
onservative 9; Mismatches 14; Indels 0; Gaps 0;

/SLGSRASLSAQEPAQEELVAEEDQDPSEINPQTEESQDPAPFLNPLVPRSPK 133
|||:|||||:|||||:|:|:|||||:|||||:|||||:|||||:|||||:|||||
/SLGSWATLSAQEPSQEELTAEDRPEPELNPQTEESQDVVPFLEQLVPRSPK 60

APRAIAAHVEVHPRFGDGAQAGVDGTVSGWEEARINSSPLRYNRIQGEFIVTRA 193
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
PRRAIAAHVEVHPRFGDGAQAGVDGTVSGWEEKINSSPLRYDRIQGEFTVIRA 120

LYCQVHFDGKAVYLLKDLLVNGVLALRCLEEFSAATAASSLGPQLRLCQVSGLLAL 253
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
LYCQVHFDGKAVYLLKDLLVNGVLALRCLEEFSAATAASSPGQLRLCQVSGLLAL 180

SLRIETLPWAHLKAAPFLTYFGLFQVH 284
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
SLRIETLPWAHLKAAPFLTYFGLFQVH 211

April 7, 2004, 17:44:47
; secs

GenCore version 5.1.6
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Search, using sw model

11 8, 2004, 19:08:51 : Search time 5584 Seconds
(without alignments)
10657.232 Million cell updates/sec

-09-245-198A-3

73
atgtcattgttagacttga.....gacaaaattgtgataaatgg 1373

GO_NUC

pop 60.0 , Gapext 60.0

70272 seqs, 21671516995 residues

ts satisfying chosen parameters: 6940544

3th: 0

3th: 2000000000

listing first 100 summaries

enEmbl:

gb_ba:

gb_hg:

gb_in:

gb_ov:

gb_ov:

gb_pat:

gb_ph:

gb_pl:

gb_pr:

gb_ro:

gb_scs:

gb_sy:

gb_un:

gb_vl:

em_ba:

em_fun:

em_hum:

em_in:

em_mu:

em_or:

em_ov:

em_pat:

em_ph:

em_pl:

em_ro:

em_scs:

em_un:

em_vl:

em_hg_hum:

em_hg_inv:

em_hg_inv:

em_hg_inv:

em_hg_inv:

em_hg_inv:

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em_hg_inv:

score greater than or equal to the score of the result being pri
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	1373	100.0	1373	6	BD062758	BD062758 A
2	1285	93.6	1306	9	AF030099	AF030099 Hoi
3	1172	85.4	1353	6	AX201324	AX201324 Se
4	1172	85.4	1353	9	AY358870	AY358870 Hoi
5	1172	85.4	1368	9	AF055872	AF055872 Hoi
6	1172	85.4	1421	6	BD090952	BD090952 Ap
7	958	69.8	1236	6	AR140407	AR140407 Se
8	958	69.8	1236	6	BD057124	BD057124 Me
9	796	58.0	1642	9	BC019047	BC019047 Hoi
10	625	45.5	898	6	AX180714	AX180714 Se
11	620	45.2	60268	9	AC016876	AC016876 Hoi
12	553	40.3	1816	9	AY081051	AY081051 Hoi
13	435	31.7	218485	2	AC127470	AC127470 Pa
14	179	13.0	195	6	AX379024	AX379024 Se
15	145	10.6	180222	2	AC130192	AC130192 Su
16	119	8.7	130254	2	AC136195	AC136195 Ra
17	119	8.7	165316	2	AC119115	AC119115 Ra
18	119	8.7	203083	2	AC069459	AC069459 Mu
19	119	8.7	203083	2	AC098923	AC098923 Ra
20	119	8.7	203083	2	AC135663	AC135663 Ra
21	119	8.7	203083	2	AL603707	AL603707 M
22	119	8.7	203083	2	AC126237	AC126237 Ca
23	119	8.7	203083	2	AF030100	AF030100 M
24	119	8.7	203083	2	AC126921	AC126921 Bo
25	119	8.7	203083	2	AC128925	AC128925 Ca
26	119	8.7	203083	2	AC126239	AC126239 Fe
27	119	8.7	203083	2	AX201395	AX201395 Se
28	119	8.7	203083	2	BD090954	BD090954 Af
29	119	8.7	203083	2	AC129071	AC129071 Pa
30	119	8.7	203083	2	BD062757	BD062757 A
31	119	8.7	203083	2	AC118309	AC118309 Ra
32	119	8.7	203083	2	AX201459	AX201459 Se
33	119	8.7	203083	2	AX522872	AX522872 Se
34	119	8.7	203083	2	AX522345	AX522345 Se
35	119	8.7	203083	2	AX522465	AX522465 Se
36	119	8.7	203083	2	AX522295	AX522295 Se
37	119	8.7	203083	2	AX209050	AX209050 Se
38	119	8.7	203083	2	BC062447	BC062447 Hc
39	119	8.7	203083	2	AR427020	AR427020 Se
40	119	8.7	203083	2	BD122573	BD122573 Ec
41	119	8.7	203083	2	AX198506	AX198506 Se
42	119	8.7	203083	2	BD122575	BD122575 Ec
43	119	8.7	203083	2	AR427022	AR427022 Se
44	119	8.7	203083	2	BC032480	BC032480 Hc
45	119	8.7	203083	2	AX330518	AX330518 Se
46	119	8.7	203083	2	AX410097	AX410097 Se
47	119	8.7	203083	2	AX895134	AX895134 Se
48	119	8.7	203083	2	BD030667	BD030667 Se
49	119	8.7	203083	2	AX381620	AX381620 Se
50	119	8.7	203083	2	BD224095	BD224095 Se
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yoda; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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nazi. A.J., Goddard, A., Godowski, P.J., Gurney, A.L.,
n.K.J., Marsters, S.A., Pan, J., Pitti, R.M., Roy, M.A., Smith, V.,
D.M., Watanabe, C.K. and Wood, W.I.
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LOCUS	AY358870	1353 bp	mRNA	
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ers,S.A., Sheridan,J.P., Pitti,R.M., Brush,J., Goddard,A. and
nazi,A., Sheridan,J.P., Pitti,R.M., Brush,J., Goddard,A. and
ification of a ligand for the death-domain-containing receptor
Biol. 8 (9), 525-528 (1998)
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ases 1 to 1368)
ers,S.A., Sheridan,J.P., Pitti,R.M., Brush,J., Goddard,A. and
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DEFINITION	Sequence 1 from patent US 6207642.		
ACCESSION	AR140407		
VERSION	AR140407.1	GI:14482903	
KEYWORDS	Unknown.		
SOURCE	Unknown.		
ORGANISM	Unknown.		
REFERENCE	1 (bases 1 to 1236)		
AUTHORS	Wiley,S.R.		
TITLE	Member of the TNF family useful for treatment and diagnosis		

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      Wiley, S.R.
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      PD 04-SEP-2001
      PF 12-FEB-1998 JP 1998535077
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REFERENCE	1		
AUTHORS	Wiley, S.R.		
TITLE	Tweak receptor		
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piens (human)

piens
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es 1 to 60268

B., Nusbaum, C. and Lander, E.

piens chromosome 17, clone RP11-186B7

shed

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Submission
 ed (08-DEC-1999) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA

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B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
 I., Bastien, V., Bloom, T., Boguslavsky, L., Bouckgalter, B.,
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TITLE JOURNAL

REFERENCE AUTHORS

Submitted (08-OCT-2002) Whitehead Institute/MIT Center for
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 4 (bases 1 to 60268)

Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Andersc
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 Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R.,
 Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schnupack,
 Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann,
 Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham,
 Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X.,
 Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zc

Direct Submission

TITLE JOURNAL

COMMENT

Submitted (31-OCT-2002) Whitehead Institute/MIT Center for
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Oct 31, 2002 this sequence version replaced gi:23592141.

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Rese

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence_submission@genome.wi.mit.edu

----- Project Information

Center project name: L3849

Center clone name: 186_B_7

FEATURES

source

Location/Qualifiers

1. .60268

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/mol_type="genomic DNA"

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/map="17"

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/clone_lib="RP11-186B7"

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/rpt_family="AluSx"

complement(8822..9119)

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14360..14438

/rpt_family="TGGn"

complement(17883..18086)

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complement(18108..18483)

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repeat_region

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repeat_region

Query Match 45.2%; Score 620; DB 9; Length 60268;
Best Local Similarity 99.6%; Pred.No. 0;
Matches 770; Conservative 0; Mismatches 3; Indels 0; C

RESULT 12			
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LOCUS	Homo sapiens TWE-PRIL mRNA, complete cds.		
DEFINITION			

1
1.1 GI:24934973
piens (human)
piens
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Primates; Catarrhini; Hominidae; Homo.
es 1 to 1816
Balade,B., Medema,J.P., Lopez-Praga,M., Lozano,J.C.,
oten,G.M., Picard,A., Martinez-A.C., Garcia-Sanz,J.A. and
genous hybrid mRNA encodes TWE-PRIL, a functional cell
TWEAK-APRIL fusion protein
21 (21), 5711-5720 (2002)
9
es 1 to 1816
Balade,B., Garcia-Sanz,J.A. and Hahne,M.
Submission
ed (21-FEB-2002) DIO, CNB, ctra de Colmenar Viejo, MADRID,
28049, Spain
Location/Qualifiers
1. .1816
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/chromosome="17"
/notes="expressed in primary T lymphocytes and monocytes"
56. .1048
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are members of the TNF ligand family, corresponding to
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/codon_start=1
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QPALRRGRGLQAGYQRIQAGVYLLYSQVLFQDVTFTMGQVVSREGQRETLFRC
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40.3%; Score 553; DB 9; Length 1816;
arity 100.0%; Pred.No. 8.3e-297; Indels 0; Gaps 0;
conservative 0; Mismatches 0; Indels 0; Gaps 0;
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CGTCGAGCCAGAGCGGAGGGGGCGGGGGGAGCGGCGACCGCCCTGCTGGT 120
CTCGGTCCTCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 230
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CTCGGTCCTCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 360
CTCGGTCCTCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 470

Db 361 GGCTGGAAGAGGATCGGACGCCCATTTAAGATTTCATCCAGACCTGGACAGGACGG
QY 471 GCAGGACAGGTGTGGACGGGACAGTGTGAGTGGTGGAGGAAGCCAGATCAACAGCTC
Db 421 GCAGGACAGGTGTGGACGGGACAGTGTGAGTGGTGGAGGAAGCCAGATCAACAGCTC
QY 531 CCCTCTGGCTACACCCGCGAGATCGGAGTTTATAGTCACCCGGGCTGGGCTCTY
Db 481 CCCTCTGGCTACACCCGCGAGATCGGAGTTTATAGTCACCCGGGCTGGGCTCTY
QY 591 CCTGTACTGTCTAG 603
Db 541 CCTGTACTGTCTAG 553
RESULT 13
AC127470 218485 bp DNA linear HTG 05:
LOCUS Pan troglodytes clone RP43-145D13, WORKING DRAFT SEQUENCE,
DEFINITION ordered pieces.
AC127470
AC127470.4 GI:31415893
VERSION HTG: HTGS PHASE2; HTGS DRAFT.
KEYWORDS Pan troglodytes (chimpanzee)
SOURCE Pan troglodytes
ORGANISM Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
REFERENCE 1 (bases 1 to 218485)
AUTHORS Antonellis,A., Ayele,K., Beckstrom-Sternberg,S.M., Benjami
Blakesley,R.W., Bouffard,G.G., Brinkley,C., Brooks,S., Car
Chu,G., Coleman,B., Coleman,H., Engle,J., Granite,S., Guan
Gupta,J., Haghighi,P., Han,J., Hansen,N., Ho,S.-L., Hu,P.,
Hurlb,B., Idol,J.R., Karlins,E., Kwong,P., Laric,P., Lee-L
Legaspi,R., Maduro,Q.D., Maduro,V.B., Paquirigan,C., Pearson,R., Portno
Maskeri,B., McDowell,J., Schandler,K., Schueler,M.G., S
Prasad,A., Reddix-Dugue,N., Schandier,K., Thomas,P.J., Teipour
Vogt,J.L., Wetherby,K.D., Wiggins,L., Young,A. and Green,E
NISC Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 218485)
Green,E.D.
Direct Submission
Submitted (17-JUL-2002) NIH Intramural Sequencing Center,
Government Circle, Gaithersburg, MD 20877, USA
3 (bases 1 to 218485)
Green,E.D.
Direct Submission
Submitted (05-JUN-2003) NIH Intramural Sequencing Center,
Government Circle, Gaithersburg, MD 20877, USA
On Jun 5, 2003 this sequence version replaced gi:26449071.
----- Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: http://www.nisc.nih.gov
Contact: nisc.zoehhri.nih.gov
----- Project Information
Center project name: cms
Center clone name: 145D13

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicat order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones, alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.
----- Summary Statistics

sequencing vector: plasmid; n/a; 100% of reads
 hemistry: Dye-terminator Big Dye; 100% of reads
 assembly program: Phrap; version 0.990319
 consensus quality: 214085 bases at least Q40
 consensus quality: 215255 bases at least Q30
 consensus quality: 216264 bases at least Q20
 insert size: 190000; agarose-fp
 insert size: 216885; sum-of-contigs
 quality coverage: 12.65x in Q20 bases; agarose-fp
 quality coverage: 11.08x in Q20 bases; sum-of-contigs

 :: This is a 'working draft' sequence. It currently
 lists of 17 contigs. Gaps between the contigs
 represented as runs of N. The order of the pieces
 believed to be correct as given, however the sizes
 the gaps between them are based on estimates that have
 rid by the submitter.

; sequence will be replaced
 the finished sequence as soon as it is available and
 the accession number will be preserved.

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1      7548: contig of 7548 bp in length
7549      7648: gap of unknown length
7649      31917: contig of 24269 bp in length
32017: gap of unknown length
32018      50433: contig of 18416 bp in length
50434      50533: gap of unknown length
50534      95274: contig of 44741 bp in length
95275      95374: gap of unknown length
95375      99988: contig of 4614 bp in length
99989      100088: gap of unknown length
10089      109873: contig of 9785 bp in length
99874      109973: gap of unknown length
99974      117619: contig of 7646 bp in length
17620      117719: gap of unknown length
17720      128625: contig of 10906 bp in length
18626      128725: gap of unknown length
28726      143521: contig of 14796 bp in length
13522      143621: gap of unknown length
43622      150832: contig of 7211 bp in length
50833      150932: gap of unknown length
50933      156073: contig of 5141 bp in length
56074      156173: gap of unknown length
187450: contig of 31277 bp in length
36174      187550: gap of unknown length
37451      191471: contig of 3921 bp in length
31472      191571: gap of unknown length
31572      207263: contig of 15692 bp in length
37264      207363: gap of unknown length
37364      209043: contig of 1680 bp in length
39044      209143: gap of unknown length
39144      217567: contig of 8424 bp in length
17568      217667: gap of unknown length
17668      218485: contig of 818 bp in length.
  
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Location/Qualifiers

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  vector_side:right"

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Best Local Similarity 99.3%; Pred. No. 1.1e-230;
Matches 755; Conservative 0; Mismatches 4; Indels 1; (
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Db 52585 CAGGTGCACTTTGATGAGGGGAAGCTGTCTACCTGAAGCTGGAGTTCCTGGTGG;
QY 661 GTGCTGGCCCTGCGCTGCTGAGGAAATTTCTCAGCACTTGGGGCCAGTTCCTCTCG;
Db 52645 GTGCTGGCCCTGCGCTGCTGAGGAAATTTCTCAGCACTTGGGGCCAGTTCCTCTCG;
QY 721 CAGCTCCCGCTCTGCGCAGGTGTCTGGGGCTGTGGCCCTGCGGCCAGGGTCTCTCC;
Db 52705 CAGCTCCCGCTCTGCGCAGGTGTCTGGGGCTGTGGCCCTGCGGCCAGGGTCTCTCC;
QY 781 ATCCGACCCCTCCCTGGGCCCATCTCAAGGTGCCCCCTTCTCAGCTACTTCTG;
Db 52765 ATCCGACCCCTCCCTGGGCCCATCTCAAGGTGCCCCCTTCTCAGCTACTTCTG;
QY 841 TTCCAGGTTCACTGAGGGGCCCTGTCTCCCACTGCTGCCAGGCTGCCGGCT;
Db 52825 TTCCAGGTTCACTGAGGGGCCCTGTCTCCCACTGCTGCCAGGCTGCCGGCT;
QY 901 CGACAGCTCTCTGGGCACCCGGTCCCTCTGCCCCCACTCCAGCCGCTCTTTGCT;
Db 52885 CGACAGCTCTCTGGGCACCCGGTCTCTCTGCCCACTCCAGCTCTTTGCT;
QY 961 CTGCCCCCTCCCTCTAGAGGCTGCTGGGCCCTGTTCAGCTGTTTCCATCCCA;
Db 52945 CTGCCCCCTCCCTCTAGAGGCTGCTGGGCCCTGTTCAGCTGTTTCCATCCCA;
QY 1021 ACAGATTCCCACTCTTATCTTACAACT - CCCCCACCGCCCACTCTCCACCTCAC;
Db 53005 ACAGATTCCCACTCTTATCTTACAACTCCCCCACTCCCACTCTCCACCTCAC;
QY 1080 CCCCATCCCTGAGCCCTTTGAGGCCCCCACTGATCTCGACTCCCCCTGGCCACA;
Db 53065 CCCCATCCCTGAGCCCTTTGAGGCCCCCACTGATCTCGACTCCCCCTGGCCACA;
QY 1140 CAGGGGCATTTGTTCACTGTACTCTGTGGGCAAGATGGTCCAGAACCCCA;
Db 53125 CCAGGGCATTTGTTTCACTGTACTCTGTGGGCAAGATGGTCCAGAACCCCA;
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Location/Qualifiers

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      8.7%; Score 119; DB 2; Length 130254;
  rity 100.0%; Pred. No. 3.9e-54; Indels 0; Gaps 0;
  nservative 0; Mismatches 0;

TCGGGTCCCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 110
|||||
TCGGGTCCCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 64163
|||||

GTCGGAGCCAGAGCGGAGGGGGCGCCGGGGGAGCGCGGACCGCCCTGCTGG 169
|||||
GTCGGAGCCAGAGCGGAGGGGGCGCCGGGGGAGCGCGGACCGCCCTGCTGG 64222
|||||

5      165316 bp      DNA      linear      HTG 19-NOV-2002
norvegicus clone CH230-320N23, WORKING DRAFT SEQUENCE, 5
ed pieces.

5.4 GI:25100662
GS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
norvegicus (Norway rat)
norvegicus
ta; Metazoa; Chordata; Vertebrata; Euteleostomi;
a; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

ies 1 to 165316)
1.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
echi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
I.D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,

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8.7%; Score 119; DB 2; Length 130254; rity 100.0%; Pred. No. 3.9e-54; Indels 0; Gaps 0; nservative 0; Mismatches 0;

TCGGGTCCCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 110
 |||||
 TCGGGTCCCGGATGGGGGGCGGTGAGGAGGACAGCCCGCCCGCCCATGGC 64163
 |||||

GTCGGAGCCAGAGCGGAGGGGGCGCCGGGGGAGCGCGGACCGCCCTGCTGG 169
 |||||
 GTCGGAGCCAGAGCGGAGGGGGCGCCGGGGGAGCGCGGACCGCCCTGCTGG 64222
 |||||

5 165316 bp DNA linear HTG 19-NOV-2002
 norvegicus clone CH230-320N23, WORKING DRAFT SEQUENCE, 5
 ed pieces.

5.4 GI:25100662
 GS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
 norvegicus (Norway rat)
 norvegicus

ta; Metazoa; Chordata; Vertebrata; Euteleostomi;
 a; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

ies 1 to 165316)
 1.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
 ., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
 echi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
 I.D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
 ., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
 N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,

TITLE

Unpublished
 2 (bases 1 to 165316)

Worley K.C.

AUTHORS

Direct Submission

JOURNAL

Submitted (25-APR-2002) Human Genome Sequencing Center, Def
 of Molecular and Human Genetics, Baylor College of Medicine
 Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 165316)

Rat Genome Sequencing Consortium.

Direct Submission

AUTHORS

Direct Submission

JOURNAL

Submitted (19-NOV-2002) Human Genome Sequencing Center, Def
 of Molecular and Human Genetics, Baylor College of Medicine
 Baylor Plaza, Houston, TX 77030, USA

On Nov 19, 2002 this sequence version replaced gi:23616728.

The sequence in this assembly is a combination of BAC bases
 and whole genome shotgun sequencing reads assembled using
 (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig de
 in the feature table below represents a scaffold in the AC
 assembly (a 'contig-scaffold'). Within each contig-scaffold
 individual sequence contigs are ordered and oriented, and
 by sized gaps filled with Ns to the estimated size. The sec
 may extend beyond the ends of the clone and there may be se
 contigs within a contig-scaffold that consist entirely of
 genome shotgun sequence reads. Both end sequences and whole
 shotgun sequence only contigs will be indicated in the feat
 table.

COMMENT

----- Genome Center
 Center: Baylor College of Medicine
 Center code: BCM

Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
 Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., C
 Cleveland,C., Cockrell,R., Cox,C., Coyie,M., Cree,A., D'Sou
 Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederic
 Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K
 Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Ea
 Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,F., Fan,G
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 Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Gar
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 Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S., Park
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 Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Rui,S., S
 Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., S
 Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.
 Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
 Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Tayl
 Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Uma
 Valas,R., Vera,V., Villaseana,D., Waldron,L., Walker,B., War
 Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F., W
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 Weinstein,G. and Gibbs,R.A.

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
 Mus musculus
 Mus musculus (house mouse)
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 Metzker, M.L., Lewis, L.R., Hume, J., Edwards, C., Harris, C., Dederich, D., Thomas, S., Okwuon, G., Carlock, C., Garner, T., Addison, S., Pace, A., Williams, G., Bonnin, D., Brooks, A., Brubay, C., Bunak, C., Burkett, C., Chacko, J., Chen, G., Chen, Z., Cox, C., Davis, C., Delgado, O., Ding, Y., Dugan-Rocha, S., Fernandez, C., Ferraguto, D., Forcum-Tansey, J., Gill, R., Gorrell, J.H., Gunaratne, P., Haller, G., Hernandez, J., Hogue Hosak, H., Hou, X., Huber, O., Jackson, L., Jia, Y., Kelly, J., Kovar, C., Liu, J., Liu, W., Loulès, H., Lozado, R.J., Morri Massey, E., McLeod, M.P., Mei, G., Moore, S., Morgan, M., Neal, D., Nelson, A., Nguyen, R., Nguyen, N., Ogih, M., Parish, Perez, L., Reiter, D., Say, J., Shen, H., Vaequez, L., Watling, Williamson, A., Wrensford, G., Zhou, X., Bouck, J., Hodgson, A., Muzny, D.M., Rives, W., Scherer, S., Sodergren, E., Weinstein, Worley, K. and Gibbs, R.
 Direct Submission
 Unpublished
 2 (bases 1 to 203083)
 Worley, K.C.
 Direct Submission
 Submitted (31-MAY-2000) Human Genome Sequencing Center, De
 of Molecular and Human Genetics, Baylor College of Medicine
 Baylor Plaza, Houston, TX 77030, USA
 On Jun 25, 2001 this sequence version replaced gi:12621364
 ----- Genome Center
 Center: Baylor College of Medicine
 Center code: BCM
 Web site: <http://www.hgsc.bcm.tmc.edu/>
 Contact: hgsc-help@bcm.tmc.edu
 Project Information
 Center project name: WAO
 Center clone name: RP23-169P5
 ----- Summary Statistics
 Sequencing vector: M13; L08821
 Chemistry: Dye-primer Bodipy: 48% of reads
 Chemistry: Dye-terminator Big Dye: 52% of reads
 Assembly program: Phrap; version 0.930329
 Consensus quality: 212648 bases at least Q40
 Consensus quality: 218902 bases at least Q30
 Consensus quality: 22384 bases at least Q20
 Estimated insert size: 210656; sum-of-contigs estimation
 Quality coverage: 0x in Q20 bases; agarose-fp estimation
 Quality coverage: 7.2x in Q20 bases; sum-of-contigs estimation

 * NOTE: Estimated insert size may differ from sequence length
 (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft.de)
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 7 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.
 * 1 6152: contig of 62152 bp in length
 * 62153 62252: gap of unknown length
 * 62253 118772: contig of 56520 bp in length
 * 118773 118872: gap of unknown length
 * 118873 148924: contig of 30052 bp in length
 * 148925 149024: gap of unknown length
 * 149025 167331: contig of 18207 bp in length
 * 167332 167331: gap of unknown length
 * 167332 189907: contig of 22576 bp in length
 * 189908 190007: gap of unknown length
 * 190008 196537: contig of 6530 bp in length
 * 196538 196637: gap of unknown length
 * 196638 203083: contig of 6446 bp in length.


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/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-154B15"
1..1822
/note="wgs contig"
complement(217607..218056)
/note="clone boundary"
clone_end:77
site:EcORI
end_sequence:RWBB0087B"

      8.7%  Score 119;  DB 2;  Length 223877;
arity 100.0%;  Pred. No. 4e-54;
conservative 0;  Mismatches 0;  Indels 0;  Gaps 0;

CTCGGTCCTCCGGATGGGGGGGGGGTGGAGGACGACAGCCCGCCCGCCCATGGC 110
|||||
CTCGGTCCTCCGGATGGGGGGGGGGTGGAGGACGACAGCCCGCCCGCCCATGGC 139398
|||||

TCGTCGAGCCAGAGCGGAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG 169
|||||
TCGTCGAGCCAGAGCGGAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG 139457
|||||

663      225077 bp      DNA      linear      HTG 10-MAY-2003
norvegicus clone CH230-46E21, WORKING DRAFT SEQUENCE.

663.3      GI:30521905
HTGS_PHASE2; HTGS_DRAFT; HTGS_FULLTOP.
norvegicus (Norway rat)
norvegicus
ota; Metazoa; Chordata; Vertebrata; Euteleostomi;
lia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
3.
ises 1 to 225077)
D.Marie, Metzker M.Lee, Abramson S., Adams C., Alder J.,
C.Allen, H. Alabrooks S., Amin A., Anguiano D.,
bechi V., Aoyagi A., Ayodeji M., Baca E., Baden H.,
in D., Bandaranaike D., Barber M., Barnstead M., Benahmed F.,
to K., Blair J., Blankenburg K., Blyth P., Brown M.,
C.N., Buhal J., Burch P., Burrell K., Calderon E.,
as V., Carter K., Cavazos I., Ceasar H., Center A.,
land C., Chavez D., Chen G., Chen R., Chen Y., Chen Z., Chu J.,
a M.L., Davis C., Davy-Carroll L., De Anda C., Dederich D.,
to O., Denson S., Deramo C., Ding Y., Dinh H., Divya K.,
r H., Dugan-Rocha S., Dunn A., Durbin K., Duval B., Eaves K.,
A., Escotto M., Eugene C., Evans C.A., Falls T., Fan G.,
ndez S., Finley M., Flagg N., Forbes L., Foster M., Foster P.,
r C.M., Gabisi A., Ganta R., Garcia A., Garner T., Garza M.,
George E., Geer K., Gill R., Grady M., Guerra W., Guevara M.,
atne P., Haaland W., Hamil C., Hamilton C., Hamilton K.,
y Y., Havlak P., Hawes A., Henderson N., Hernandez J.,
ndez R., Hines S., Hladun S.L., Hodgson A., Hogues M.,
ns B., Howells S., Hulyk S., Hume J., Idlebird D., Jackson A.,
on L., Jacob L., Jiang H., Johnson B., Johnson R., Jolivet A.,
thy S., Kelly S., Kelly S., Khan Z., King L., Kovar C.,
C., Kraft C.L., Lebow H., Levan J., Lewis L., Li Z., Liu J.,
shewa L., Louisgied H., Lozada R.J., Lu X., Ma J.,
hwari M., Mahindartine M., Mahmoud M., Malloy K., Mangum A.,
n B., Mapua P., Martin K., Martin R., Martinez E.,
ney S., McLeod M.P., McNeill T.Z., Meenen E.,
aviljevic A., Miher G., Minja E., Montemayor J., Moore S.,
n M., Morris K., Morris S., Munidasa M., Murphy M., Nair L.,
elvis C., Neal D., Newton N., Nguyen N., Norris S.,
elenah O., Okwuonu G., Olarnpunsagoon A., Pal S., Parks K.,
rnak S., Paul H., Perez A., Perez L., Pfannkoch C.,
er F., Poindexter A., Popovic D., Primus E., Pu L.-L.,
M., Quiroz J., Rachlin E., Reeves K., Regier M.A., Reigh R.,

```

```

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

1 (bases 1 to 225077)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (06-NOV-2002) Human Genome Sequencing Center, De
of Molecular and Human Genetics, Baylor College of Medicin
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 225077)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (10-MAY-2003) Human Genome Sequencing Center, De
of Molecular and Human Genetics, Baylor College of Medicin
Baylor Plaza, Houston, TX 77030, USA
The sequence in this assembly is a combination of BAC base
and whole genome shotgun sequencing reads assembled using
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig (
in the feature table below represents a scaffold in the A
assembly (a 'contig-scaffold'). Within each contig-scaffo
individual sequence contigs are ordered and oriented, and
by sized gaps filled with Ns to the estimated size. The s
may extend beyond the ends of the clone and there may be
contigs within a contig-scaffold that consist entirely of
genome shotgun sequence reads. Both end sequences and who
shotgun sequence only contigs will be indicated in the fe
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GHDQ
Center clone name: CH230-46E21
----- Summary Statistics
Assembly program: Atlas 3.0;
Consensus quality: 214599 bases at least Q40
Consensus quality: 216978 bases at least Q30
Consensus quality: 218480 bases at least Q20
Estimated insert size: 225046; sum-of-contigs estimat
Quality coverage: 8x in Q20 bases; sum-of-contigs est

* NOTE: Estimated insert size may differ from sequence le
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_d
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that ha
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 225077: contig of 225077 bp in length.
1. 225077 Location/Qualifiers
/organism="Rattus norvegicus"
/mol_type="genomic DNA"

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FEATURES
source

```

/db_xref="taxon:10116"
/clone="CH230-46E21"
1. .1402
/note="wgs_contig"

      8.7%; Score 119; DB 2; Length 225077;
urity 100.0%; Pred. NO. 4e-54;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

TTCGTCCTCCGGATCGGGGGGGCGGTGAGCAGGACACAGCCCGCCCGCCATGGC 110
TTCGTCCTCCGGATCGGGGGGGCGGTGAGCAGGACACAGCCCGCCCGCCATGGC 199986

TGTCCGAGCAGAGCGAGGGGGCGCGGGGGAGCCGGGACCCCGCTGTGG 169
TGTCCGAGCAGAGCGAGGGGGCGCGGGGGAGCCGGGACCCCGCTGTGG 199927

07      234182 bp DNA linear ROD 17-NOV-2001
DNA sequence from clone RP23-422L16 on chromosome 11,
ie sequence.
07.5 GI:17017790
iculus (house mouse)
ta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
a; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
ses 1 to 234182)
A.
Submission
ed (17-NOV-2001) Wellcome Trust Sanger Institute, Hinxton,
ighshire, CB10 1SA, UK. E-mail enquiries:
ysanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
20, 2001 this sequence version replaced gi:16605765.
sequence assembly data is compared from overlapping clones.
ifferences are found these are annotated as variations
on with a note of the overlapping clone name. Note that the
ion annotation may not be found in the sequence submission
onding to the overlapping clone, as we submit sequences with
small overlap as described above.
sequence was finished as follows unless otherwise noted: all
s were either double-stranded or sequenced with an alternate
try or covered by high quality data (i.e., phred quality >=
1 attempt was made to resolve all sequencing problems, such
pressions and repeats; all regions were covered by at least
amid subclone or more than one M13 subclone; and the
ly was confirmed by restriction digest. The following
iations are used to associate primary accession numbers given
feature table with their source databases: Em, EMBL; Sw,
ROT; Tr, TREMBL; Wp, WORMPEP; information on the WORMPEP
se can be found at
/www.sanger.ac.uk/Projects/C_elegans/wormpep RP23-422L16 is
ne RPCI-23 Mouse PAC Library
acted by the group of Pieter de Jong.
rther details see http://www.chori.org/bacpac/home.htm
: pBACe3.6
Location/Qualifiers
1. .234182
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
/chromosome="11"
/clone="RP23-422L16"
/clone_lib="RPCI-23"
complement(84050..84131)
/note="Sequence from uni-directional primer reads and dGTP
big dye terminator reads only."

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Query Match      8.7%; Score 119; DB 10; Length 234182;
Best Local Similarity 100.0%; Pred. No. 4.1e-54;
Matches 119; Conservative 0; Mismatches 0; Indels 0; G:

QY      51 ATCCCTCCGGATCGGGGGGGCGGTGAGCAGGACACAGCCCGCCCGCCATGGC
DB      75038 ATCCCTCCGGATCGGGGGGGCGGTGAGCAGGACACAGCCCGCCCGCCATGGC
QY      111 CGCCCTCCGAGCAGAGCGGAGGGGGCGCGGGGGAGCCGGGACCCCGCTGTGC
DB      74978 CGCCCTCCGAGCAGAGCGGAGGGGGCGCGGGGGAGCCGGGACCCCGCTGTGC

RESULT 22
AC126237      212093 bp DNA linear HTG 06
LOCUS
DEFINITION
Canis familiaris clone RP81-414022, WORKING DRAFT SEQUENCE
ordered pieces.
AC126237
AC126237.5 GI:31442445
VERSION
HTG; HTGS PHASE2; HTGS DRAFT.
KEYWORDS
SOURCE
Canis familiaris (dog)
ORGANISM
Canis familiaris
REFERENCE
1 (bases 1 to 212093)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele-
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
Antonellis,A., Ayele,K., Beckstrom-Sternberg,S.M., Benjami
Blakesley,R.W., Bouffard,G.G., Brinkley,C., Brooks,S., Car
Chu,C., Coleman,B., Coleman,H., Engle,J., Granite,S., Guan
Gupta,J., Haghighi,P., Han,J., Hansen,N., Ho,S.-L., Hu,P.,
Hurtle,B., Idol,J.R., Karlins,E., Kwong,P., Laric,P., Lee-L
Legaspi,R., Maduro,Q.L., Maduro,V.B., Margulies,E.H., Masi
Maskeri,B., McDowell,J., Paquirigan,C., Pearson,R., Porto
Prasad,A., Reddix-Dugue,N., Schandler,K., Schueler,M.G., S
Sison,C., Stantropop,S., Thomas,J.W., Thomas,P.J., Tsipour
Vogt,J.L., Wetherby,K.D., Wiggins,L., Young,A. and Green,E
NISC Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 212093)
Green,E.D.
Direct Submission
Submitted (04-JUL-2002) NIH Intramural Sequencing Center,
Grovemont Circle, Gaithersburg, MD 20877, USA
3 (bases 1 to 212093)
Green,E.D.
Direct Submission
Submitted (06-JUN-2003) NIH Intramural Sequencing Center,
Grovemont Circle, Gaithersburg, MD 20877, USA
On Jun 6, 2003 this sequence version replaced gi:27476131.
----- Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: http://www.nisc.nih.gov
Contact: nisc_zoo@ngri.nih.gov
----- Project Information
Center project name: cwq
Center clone name: 414022

```

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicat order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones, alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

----- Summary Statistics -----
Sequencing vector: plasmid; n/a; 100% of reads

```

/notes="clone overlaps with GenBank Accession Number
AC126925 clone RP81-332E11 (center project name cwp)"
137892. .146862

```



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/note="assembly_fragment"
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/note="clone overlaps with GenBank Accession Number
AC134961 clone RP42-406J16 (center project name djv)"
126979..129099
/note="assembly_fragment"
129200..135799
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135900..145102
/note="assembly_fragment"
145203..148555
/note="assembly_fragment
clone_end:77
vector_side:right"

4.2%; Score 58; DB 2; Length 148555;
larity 100.0%; Pred. No. 4.7e-20;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GTGACCTTGTGAGGGAGGCTGCTACCTGAAGCTGGACTGCTGTGGATG 658
|||||
GTGACCTTGTGAGGGAGGCTGCTACCTGAAGCTGGACTGCTGTGGATG 60929

925 176258 bp DNA linear HTG 06-JUN-2003
familiaris clone RP81-332E11, WORKING DRAFT SEQUENCE, 12
ed pieces.
925.6 GI:31442444
HTGS PHASE2; HTGS_DRAFT.
familiaris (dog)
familiaris
yota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
lia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
ases 1 to 176258)
ellis A., Ayele K., Beckstrom-Sternberg S.M., Benjamin B.,
sley R.W., Bouffard G.G., Brinkley C., Brooks S., Cariaga K.,
Coleman B., Coleman H., Engle J., Hansen N., Ho S.-L., Hu P.,
J., Haghighi P., Han J., Han J., Laric P., Lee-Lin S.-Q.,
B., Idol J.R., Karlins E., Kwong P., Margulies E.H., Mastello C.,
pi R., Maduro Q.L., Maduro V.B., Margulies E.H., Mastello C.,
ri B., McDowell J., Paquinigan C., Pearson R., Portnoy M.E.,
d A., Reddix-Dugue N., Schandler K., Schueler M.G., Shah X.,
C., Stantropop S., Thomas J.W., Thomas P.J., Tsipouri V.,
J.L., Wetherby K.D., Wiggins L., Young A. and Green E.D.
Comparative Sequencing Initiative
lished
ases 1 to 176258)

t Submission
tted (10-JUL-2002) NIH Intramural Sequencing Center, 8717
mont Circle, Gaithersburg, MD 20877, USA
ases 1 to 176258)
E.D.

t Submission
tted (06-JUN-2003) NIH Intramural Sequencing Center, 8717
mont Circle, Gaithersburg, MD 20877, USA
n 6, 2003 this sequence version replaced gi:28209436.
----- Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: http://www.nisc.nih.gov
Contact: nisc.zoo@nih.gov
----- Project Information
Center project name: CWP
Center clone name: 332E11

```

sequence data in this record represents an 'enhanced' on of a Phase 2 submission. Specifically, the indicated and orientation of each sequence contig has been

established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones, alignment with available reference sequence (e.g. human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

----- Summary Statistics

Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 173760 bases at least Q40
Consensus quality: 174423 bases at least Q30
Consensus quality: 174916 bases at least Q20
Insert size: 152000; agarose-fp
Insert size: 175158; sum-of-contigs
Quality coverage: 17.46x in Q20 bases; agarose-fp
Quality coverage: 15.15x in Q20 bases; sum-of-contig

* NOTE: This is a 'working draft' sequence. It currently
* consists of 12 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that ha
* provided by the submitter.

* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

```

1 11425: contig of 11425 bp in length
* 11426 11525: gap of unknown length
* 11526 27554: contig of 16029 bp in length
* 27555 29974: gap of unknown length
* 29975 30074: contig of 2320 bp in length
* 30075 36241: gap of unknown length
* 36242 36342: contig of 6167 bp in length
* 36343 77002: contig of 40661 bp in length
* 77003 77103: gap of unknown length
* 77104 117018: contig of 39916 bp in length
* 117019 117119: gap of unknown length
* 117120 119041: contig of 1923 bp in length
* 119042 119141: gap of unknown length
* 119142 158388: contig of 39247 bp in length
* 158389 158489: gap of unknown length
* 158490 168033: contig of 9545 bp in length
* 168034 168133: gap of unknown length
* 168134 170716: contig of 2583 bp in length
* 170717 170816: gap of unknown length
* 170817 174429: contig of 3613 bp in length
* 174430 174529: gap of unknown length
* 174530 176258: contig of 1729 bp in length.

```

FEATURES

	Location/Qualifiers	source
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		/mol_type="genomic DNA"
		/db_xref="taxon:9615"
		/clone="RP81-332E11"
		/clone_lib="RP81"
misc_feature	1..88033	/note="clone overlaps with GenBank Accession Num AC126237 clone RP81-414022 (center project name
misc_feature	1..11425	/note="assembly_fragment clone_end:896 vector_side:left"
misc_feature	11526..17554	/note="assembly_fragment"
misc_feature	27655..29974	/note="assembly_fragment"
misc_feature	30075..36241	/note="assembly_fragment"

/note="assembly_fragment"
24417. .36979
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37080. .42619
/note="assembly_fragment"
42720. .46464
/note="assembly_fragment"
46565. .61551
/note="assembly_fragment"
61652. .92225
/note="assembly_fragment"
93226. .110459
/note="assembly_fragment"
110560. .147587
/note="assembly_fragment"
147688. .149736
/note="assembly_fragment"
clone_end:SP6
vector_side:right"

3.9%; Score 53; DB 2; Length 149736;
Identity 100.0%; Pred. No. 3e-17; 0; Indels 0; Gaps 0;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CAGGGTCTCTCCCTGGGATCCGACCCCTCCCTGGGCGCATCTCAAGGC 812
CAGGGTCTCTCCCTGGGATCCGACCCCTCCCTGGGCGCATCTCAAGGC 43768

395 ace 74 from Patent WO0153486. linear PAT 30-AUG-2001
395

395.1 GI:15391196

atic construct
etic construct
lcial sequences.

razi,A.J., Goddard,A., Godowski,P.J., Gurney,A.L.,
1,K.J., Marsters,S.A., Pan,J., Pitti,R.W., Roy,M.A., Smith,V.,
D.M., Watanabe,C.K. and Wood,W.I.
sitions and methods for the treatment of tumour
-: WO 0153486-A 74 26-JUL-2001;
tech, Inc. (US)

Location/Qualifiers

1. .50
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic Oligonucleotide Probe."

3.6%; Score 50; DB 6; Length 50;
Identity 100.0%; Pred. No. 7e-16;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GCCTCTCGCTACACCGCAGATCGGGAGTTTATAGTCACCCGG 576
GCCTCTCGCTACACCGCAGATCGGGAGTTTATAGTCACCCGG 50

954 ligand polypeptide. linear PAT 27-AUG-2002
954

954.1 GI:22636564

01522584-A/3.

ntified
ntified

unclassified

1 (bases 1 to 50)
Askenazi,A.J., Marsters,S.A. and Pitti,R.
Apo-3 ligand polypeptide
Patent: JP 2001522584-A 3 20-NOV-2001;
GENENTECH INC

OS Unknown
PN JP 2001522584-A/3

PD 20-NOV-2001
PF 09-OCT-1998 JP 2000516042
PR 10-OCT-1997 US 60/062037,17-DEC-1997 US 60/0691
AVI J ASHENAZI, SCOT A MARSTERS, ROBERT PITTI
PC C12N15/09,A61K38/00,C07K14/705,C07K16/24,C12N15/00,A6
CC Description of Unknown Organism:Unknown
FH Key Location/Qualifiers
FT source 1. .50
/organism="Unknown".

Location/Qualifiers

1. .50
/organism="unidentified"
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/db_xref="taxon:32644"

ORIGIN

Query Match 3.6%; Score 50; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 7e-16;
Matches 50; Conservative 0; Mismatches 0; Indels 0;

QY 527 CCAGCCCTCTCGCTACACCGCAGATCGGGAGTTTATAGTCACCCGG 576
|||||
DB 1 CCAGCCCTCTCGCTACACCGCAGATCGGGAGTTTATAGTCACCCGG 50

RESULT 29

AC129071/c

LOCUS

DEFINITION

AC129071

AC129071.2 GI:26449072

HTG; HTGS PHASE2; HTGS DRAFT.

SOURCE

ORGANISM

Pan troglodytes

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutel

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.

1 (bases 1 to 163542)

REFERENCE

AUTHORS

AKHTER,N., Antonellis,A., Ayele,K., Beckstrom-Sternberg,S

Benjamin,B., Blakesley,R.W., Bouffard,G.G., Brinkley,C.,

Benjamin,B., Coleman,B., Engle,J., Granite,S., Guan,X., Gu

Haghighi,P., Han,J., Hansen,N., Ho,S.-L., Idol,J.R., Karl

Laric,P., Lee-Lin,S.-Q., Legaspi,R., Maduro,Q.L., Maduro,

Margulies,E.H., Masello,C., Maskeri,B., McDowell,J.,

Paguirigan,C., Pearson,R., Portnoy,M.E., Prasad,A.,

Reddix-Dugue,N., Schandler,K., Schueler,M.G., Sison,C.,

Stantropop,S., Thomas,J.W., Thomas,P.J., Touchman,J.W., V

Wetherby,K.D., Wiggins,L., Young,A. and Green,E.D.

NISC Comparative Sequencing Initiative

Unpublished

2 (bases 1 to 163542)

Green,E.D.

Direct Submission

Submitted (25-JUL-2002) NIH Intramural Sequencing Center,

Groveton Circle, Gaithersburg, MD 20877, USA

3 (bases 1 to 163542)

Green,E.D.

Direct Submission

Submitted (11-DEC-2002) NIH Intramural Sequencing Center,

Groveton Circle, Gaithersburg, MD 20877, USA

On Dec 11, 2002 this sequence version replaced gi:2195501

----- Genome Center

Center: NIH Intramural Sequencing Center

Center code: NISC

Web site: http://www.nisc.nih.gov

ntact: misc.zoemhgri.nih.gov
----- Project Information
nter project name: cmt
nter clone name: 149M23

quence data in this record represents an 'enhanced'
1 of a Phase 2 submission. Specifically, the indicated
and orientation of each sequence contig has been
ished using one or more of the following: read-pair
com individual subclones, overlaps with neighboring
alignment with available reference sequence (e.g.,
and/or confirmation by PCR testing. In addition,
quence assembly is based on at least 8x average
ye in Q20 bases and has been reviewed to rule out
misassemblies, the low-quality ends of sequence
s have been trimmed away, and each base is associated
Phrap-derived quality score.

----- Summary Statistics
sequencing vector: plasmid; n/a; 100% of reads
chemistry: Dye-terminator Big Dye; 100% of reads
assembly program: Phrap; version 0.990319
consensus quality: 159675 bases at least Q40
consensus quality: 160848 bases at least Q30
insert size: 185000; agarose-fp
insert size: 162442; sum-of-contigs
ality coverage: 7.43x in Q20 bases; agarose-fp
ality coverage: 8.47x in Q20 bases; sum-of-contigs

: This is a 'working draft' sequence. It currently
ists of 12 contigs. Gaps between the contigs
represented as runs of N. The order of the pieces
elieved to be correct as given, however the sizes
he gaps between them are based on estimates that have
ided by the submittor.

sequence will be replaced
he finished sequence as soon as it is available and
accession number will be preserved.

1 10517: contig of 10517 bp in length
0518 10617: gap of unknown length
0618 25192: contig of 14575 bp in length
5193 25292: gap of unknown length
5293 27192: contig of 1900 bp in length
7193 27292: gap of unknown length
7293 29759: contig of 2467 bp in length
9760 29859: gap of unknown length
9860 57864: contig of 28005 bp in length
7865 57964: gap of unknown length
7965 61287: contig of 3323 bp in length
1288 61387: gap of unknown length
1388 84039: contig of 22652 bp in length
4040 84139: gap of unknown length
4140 114016: contig of 29877 bp in length
4017 114116: gap of unknown length
4117 116474: contig of 2358 bp in length
6475 116574: gap of unknown length
6575 121369: contig of 4795 bp in length
1370 121469: gap of unknown length
1470 154199: contig of 32730 bp in length
4200 154299: gap of unknown length
4300 163542: contig of 9243 bp in length.

Location/Qualifiers
1..163542
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="RP43-149M23"
/clone_lib="RP43"
1..10517
/note="assembly_fragment
missing T7 clone end on 5' end of insert"
10618..25192
/note="assembly_fragment"

misc_feature 25293..27192
/note="assembly_fragment"
misc_feature 27293..29759
/note="assembly_fragment"
misc_feature 29860..57864
/note="assembly_fragment"
misc_feature 57965..61287
/note="assembly_fragment"
misc_feature 61388..84039
/note="assembly_fragment"
misc_feature 84140..114016
/note="assembly_fragment"
misc_feature 114117..116474
/note="assembly_fragment"
misc_feature 116575..121369
/note="assembly_fragment"
misc_feature 121470..154199
/note="assembly_fragment"
misc_feature 154300..163542
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clone end:SP6
vector_side:right"

ORIGIN

Query Match 3.5%; Score 48; DB 2; Length 163542;
Best Local Similarity 100.0%; Pred. No. 1.9e-14;
Matches 46; Conservative 0; Mismatches 0; Indels 0; G

QY 1312 CCTGTGGATTTTAAACAGATATTATTTATTATTATTGTGACAAA 1359
Db 163542 CCTGTGGATTTTAAACAGATATTATTTATTATTATTGTGACAAA 163495

RESULT 30
BD062757 1168 bp DNA linear PAT 27
LOCUS
DEFINITION A tumor necrosis factor related ligand.
ACCESSION BD062757
VERSION BD062757.1 GI:22608360
KEYWORDS JP 2001505407-A/1.
SOURCE unidentified
ORGANISM unidentified
unclassified.

REFERENCE 1 (bases 1 to 1168)
AUTHORS Chicheportiche,Y. and Browning,J.L.
TITLE A tumor necrosis factor related ligand
JOURNAL Patent: JP 2001505407-A 1 24-APR-2001;
BIOGEN INC,THE FACULTY OF MEDICINE OF THE UNIVERSITY OF GE
COMMENT OS TNF family related protein
PN JP 2001505407-A/1
PD 24-APR-2001
PR 07-AUG-1997 JP 1998508239
PF 07-AUG-1996 US 60/023541,18-OCT-1996 US 60/0285
18-MAR-1997 US 60/040820
PI YVES CHICHEPORTICHE,JEFFREY L BROWNING
PC C12N15/28,C07K14/525,G01N33/68,C07K16/24,C12N15/11,A6
PC C12N5/10,
PC A61K39/395,A61K38/19,C07K14/705,C12N15/12
CC Strandedness: Double;
CC Topology: Linear;
FH Key Location/Qualifiers
FT CDS
Location/Qualifiers
1..1168
/organism="unidentified"
/mol_type="genomic DNA"
/db_xref="taxon:32644"

FEATURES

source
1..1168
/organism="unidentified"
/mol_type="genomic DNA"
/db_xref="taxon:32644"

ORIGIN

Query Match 3.4%; Score 46; DB 6; Length 1168;
Best Local Similarity 100.0%; Pred. No. 1.6e-13;
Matches 46; Conservative 0; Mismatches 0; Indels 0; C

Submitted (15-APR-2002) Human Genome Sequencing Center, Baylor College of Medicine, 6401 Fannin Street, Houston, TX 77030, USA

Direct Submission
Submitted (13-NOV-2002) Human Genome Sequencing Center, Dept. of Molecular and Human Genetics, Baylor College of Medicine
Baylor Plaza, Houston, TX 77030, USA

The sequence in this assembly is a combination of BAC bases and whole genome shotgun sequencing reads assembled using (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig in the feature table below represents a scaffold in the Assembly. A 'contig-scaffold' within each contig-scaffold

assembly (a 'contig-scaffold'). Within each contig-scaffold, the individual sequence contigs are ordered and oriented, and sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be contigs within a contig-scaffold that consist entirely of shotgun sequence reads. Both end sequences and whole shotgun sequence only contigs will be indicated in the following table.

contigs within a contig-scaffold that consist entirely of genome shotgun sequence reads. Both end sequences and shotgun sequence only contigs will be indicated in the following table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hpsc-hel@bcm.tmc.edu

- * as soon as it is available and the accession number will
- * be preserved.

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feature
225726. .227372
/note="wgs_end_extension
clone end.T7"
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/note="wgs end _extension
clone end.T7"

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clone_end:T7"
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ORGANISM	REFERENCE	AUTHORS	TITLE	JOURNAL	MEDLINE	PUBMED	REFERENCE	AUTHORS	TITLE	JOURNAL	REMARK	COMMENT	
Homo sapiens	1 (bases 1 to 353)	Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klauener, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schi Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bha Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsie Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, S., Stapleton, M., Soares, M.B., Donald, M.F., Casavant, T.L., Scheetz, T.E., Brownstein, M.J., Ustin, T.B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peter: Abramson, R.D., Mullaly, S.J., Bosak, S.A., McEwan, P.J., McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S.S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R. Fahey, J., Helton, E., Kettman, M., Madan, A., Rodrigues, S., Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D., Dicksen, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myi Butterfield, Y.S., Krzywinski, K.I., Skalska, U., Smallwood, D.J., Scherch, A., Schein, J.E., Jones, S.J. and Marra, M.A.	Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences	Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)	22388257		2	(bases 1 to 353)	Strausberg, R.	Direct Submission	Submitted (17-NOV-2003) National Institutes of Health, Ma Gene Collection (MGC), Cancer Genomics Office, National C Institute, 31 Center Drive, Room 11A03, Bethesda, MD 2089 USA	NIH-MGC Project URL: http://mgc.nci.nih.gov	Contact: MGC help desk Email: cgaps-remail.nih.gov Tissue Procurement: ATCC/DCTP cDNA Library Preparation: CLONTECH Laboratories, Inc. cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Genome Sequence Centre, BC Cancer Agency, Vancouver, BC, Canada info@bcgsc.bc.ca Steven Jones, Jennifer Aano, Ian Bosdet, Yaron Butterfie Susanna Chan, Readman Chiu, Chris Fjell, Erin Garland, Ra Lettigia Hsiao, Martin Krzywinski, Reta Kutsche, Oliver L Sen Lee, Victor Ling, Carrie Mathewson, Candice McEavey, J News, Pawan Pandoh, Anna-Lilaea Prabhu, Parvaneh Saeedi, J Schein, Duane Smailus, Michael Smith, Lorraine Spence, J Michael Thorne, Miranada Tsai, Natasja van den Bosch, Jil George Yang, Scott Zuyderduyn, Marco Marra.
FEATURES	source	Location/Qualifiers	1. 353										
		/organism="Homo sapiens"											
		/mol_type="mRNA"											
		/db_xref="taxon:9606"											
		/clone="IMAGE:3932215"											
		/tissue_type="Skin, melanotic melanoma, high MDR"											
		/clone_lib="NIH MGC_62"											
		/lab_host="DH10B"											
		/note="Vector: pDNR-LIB"											
ORIGIN													
Query Match		1.9%;	Score 26;	DB 9;	Length 353;								
Best Local Similarity		100.0%;	Prod. No. 0.021;										
Matches		26;	Conservative	0;	Mismatches	0;	Indels	0;					

d encoded human protein.

75.1 GI:23217520

2010789-A/14652.

apiens (human)

ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

is, J.B.D.M., Jobert, S. and Giordano, J.E.

id encoded human protein

IP 2002010789-A 14652 15-JAN-2002;

1 CORP

homo sapiens (human)

IP 2002010789-A/14652

15-JAN-2002

07-AUG-2000 JP 2000280989

05-AUG-1999 US 60/147499

JEAN BAPTISTE DUMAS MILNE EDWARDS, SEVELIN JOBERT, JEAN EVE PI

DANO

12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, PC

1/21, C12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12N5/00, PC

5/00

IST and encoded human protein

key Location/Qualifiers

source 1.416 /organism='Homo sapiens (human)'

Location/Qualifiers

1.416

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

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Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

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TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

TCATGTTAGACTTTGAAATTC 26

TCATGTTAGACTTTGAAATTC 356

Akhter, N., Ayele, K., Becketrom-Sternberg, S.M., Benjamin, B.,
Blakesley, R.W., Bouffard, G.G., Breen, K., Brinkley, C., Broc,
Dietrich, N.L., Granite, S., Guan, X., Gupta, J., Haghighi, P.,
Hansen, N., Ho, S.-L., Karlins, E., Kwong, P., Laric, P., Legas,
Maduro, Q.L., Masiello, C., Maskeri, B., Mastrian, S.D., McClos,
McDowell, J., Pearson, R., Stantropop, S., Thomas, P.J., Touchi,
Tsurgon, C., Vogt, J.L., Walker, M.A., Wetherby, K.D., Wiggir
Young, A., Zhang, L.-H. and Green, E.D.

Clone distribution: MGC clone distribution information can
through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Series: IRAK plate: 64 Row: 1 Column: 24
This clone was selected for full length sequencing because
passed the following selection criteria: Hexamer frequency
analysis.

FEATURES

source

Location/Qualifiers

1..418

/organism="Homo sapiens"

/mol_type="RNA"

/db_xref="taxon:9606"

/clone="IMAGE:5214272"

/tissue_type="Blood, adult leukocytes"

/clone_lib="NIH MGC_118"

/lab_host="DH10B"

/note="Vector: pCMV-SPORT6"

1..281

/codon_start=3

/product="Unknown (protein for IMAGE:5214272)"

/protein_id="AAH32480.1"

/db_xref="GI:21619103"

/translation="RSVLLLVAVRLHLLSCPLQBPAGTEWILEEGV
DIYNLRSPPENWNRGALWKEKDRPCAFMKVKIWLNFHKTIVVIA

ORIGIN

Query Match 1.9%; Score 26; DB 9; Length 418;

Best Local Similarity 100.0%; Pred. No. 0.022;

Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 1 ATGTCATTGTTAGACTTTGAAATTC 26

|||||

Db 309 ATGTCATTGTTAGACTTTGAAATTC 334

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ATTGTTAGACTTTGAAATTC 47

17 439 bp DNA linear PAT 14-JUN-2002
e 2744 from Patent WO0229103.
17
17.1 GI:21442802

apiens (human)
ata, Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
3,C., Horne, D., Peres-da-Silva, S. and Vockley, J.G.
pression profiles in liver cancer
e WO 0229103-A 2744 11-APR-2002;
GIC INC (US)
Location/Qualifiers
1. .439
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"
/note="EMBL/GenBank Accession No. N98464"

arity 1.9%; Score 26; DB 6; Length 439;
100.0%; Pred. No. 0.022;
conservative 0; Mismatches 0; Indels 0; Gaps 0;

CATTGTTAGACTTTGAAATTC 26
|||||
CATTGTTAGACTTTGAAATTC 47

34 452 bp DNA linear PAT 18-DEC-2003
ce 10997 from Patent EP1033401.
34
34.1 GI:40050018

apiens (human)
apiens
ota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

Milne Edwards, J.B., Duclert, A. and Giordano, J.Y.
sed sequence tags and encoded human proteins
e EP 1033401-A 10997 06-SEP-2000;
(FR)

Location/Qualifiers
1. .452
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

arity 1.9%; Score 26; DB 6; Length 452;
100.0%; Pred. No. 0.022; 0; Indels 0; Gaps 0;
conservative 0; Mismatches 0; Indels 0;

TCATTGTTAGACTTTGAAATTC 26
|||||
TCATTGTTAGACTTTGAAATTC 439

567 452 bp DNA linear PAT 27-AUG-2002
nce tag and encoded human protein.

ACCESSION BD030667
VERSION BD030667.1 GI:22572409
KEYWORDS JP 2001269182-A/6913.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 452)
AUTHORS Edwards, J.B.D.M., Duclair, E. and Jordan, J.Y.
TITLE Sequence tag and encoded human protein
JOURNAL Patent: JP 2001269182-A 6913 02-OCT-2001;
GENSET

COMMENT
OS Homo sapiens (human)
PN JP 2001269182-A/6913
PD 02-OCT-2001
PF 24-FEB-2000 JP 2000118773
PR 26-FEB-1999 US 60/122487
PI JEAN BAPTISTE DUMAS MILNE EDWARDS, EIMERIC DUCLAIR, JEA
PI JORDAN
PC C12N15/09, C07K14/435, C07K16/18, C12N1/15, C12N1/19, C12N
C12N5/10,
PC C12P21/02, C12P21/08, C12Q1/68//G06F17/30, C12N15/90, C12
G06F15/40

CC Key Location/Qualifiers.

FEATURES
source
1. .452
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 0.022;
Matches 26; Conservative 0; Mismatches 0; Indels 0; C

Qy 1 ATGTCATTGTTAGACTTTGAAATTC 26
|||||
Db 414 ATGTCATTGTTAGACTTTGAAATTC 439

RESULT 49
AX381620 483 bp DNA linear PAT 18
LOCUS Sequence 558 from Patent WO0212280.
DEFINITION AX381620
ACCESSION AX381620
VERSION AX381620.1 GI:19576442
KEYWORDS

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eutele;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo

REFERENCE 1
AUTHORS Pyle, R.A., Xu, J. and Secrist, H.
TITLE Compositions and methods for the therapy and diagnosis of
cancer
JOURNAL Patent: WO 0212280-A 558 14-FEB-2002;
CORIXA CORPORATION (US)

FEATURES
source
1. .483
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 0.022; 0; Indels 0;
Matches 26; Conservative 0; Mismatches 0;

Qy 1 ATGTCATTGTTAGACTTTGAAATTC 26
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Db 397 ATGTCATTGTTAGACTTTGAAATTC 422

iard; DNA; 153 BP.
 (first entry)
 ell derived DNA fragment #168.
 ell; immunosuppressive; immunostimulant; antiinflammatory;
 ene therapy; vaccine; allergen; transplant rejection;
 host disease; malignant disease; ds.
 2000DE-01021834.
 2000DE-01021834.
 THERAPEUTICS GMBH.
 inter H. Reinartz J;
 320/04.
 cative of T cell activation and functional status, useful
 : and therapy e.g. of autoimmunity or transplant rejection.
 : 48; 94pp; German.
 : represents a novel messenger RNA, (mRNA), (I), for use as
 the activation and functional status of T cells, that have
 reduced expression, and are present at higher or lower
 4, in activated T cells, relative to normal or resting cells,
 ridizes to any of 334 sequences, reproduced, or their
 complements or fragments. The products of the invention have
 sive, immunostimulant, antiinflammatory and cytostatic
 can be used for gene therapy. The polynucleotides of the
 : used: (i) as reagent for detecting activation/functional
 cells, for diagnosis, therapy, modulation or control of the
 ses of (auto)immunity (against microorganisms, vaccines or
 ransplant rejection; immunologically-related inflammation;
 sion; immune deficiency; guest versus host disease, and
 rmaceuticals, that bind to (iii) or derived polypeptides
 to prepare kits for measuring gene expression profiles in
 ne, especially T cells; (iv) to raise antibodies (Ab)
 net (iii); and (v) to prepare binding molecules (iv)
 (ii). Ab and (iv) are also useful for detecting and
 e activation and functional status of T cells. AAI68865-
 resent the activated T-cell derived polynucleotide fragments
 the method of the invention
 BP; 43 A; 23 C; 28 G; 59 T; 0 U; 0 Other;
 .arity 1.9%; Score 26; DB 6; Length 153;
 .arity 100.0%; Pred. No. 0.11;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 TCATTGTTAGACTTTGAAATTC 26
 |||||
 TCATTGTTAGACTTTGAAATTC 55

DT 07-NOV-2001 (first entry)
 XX Human ovarian PCR-subtracted cDNA library clone #890.
 DE Immunogenic protein; cancer; ovarian tumour; T-cell stimulation;
 XX gene therapy; cytostatic; T-cell expansion; nucleic acid hybridis
 KW primer; probe.
 KW Homo sapiens.
 XX WO200157207-A2.
 PN 09-AUG-2001.
 XX 05-FEB-2001; 2001WO-US003733.
 XX 04-FEB-2000; 2000US-0180403P.
 PR 28-MAR-2000; 2000US-0192745P.
 XX (CORI-) CORIXA CORP.
 XX Algate PA, Mannion J;
 PI WPI; 2001-488879/53.
 XX New polynucleotides encoding ovarian tumor proteins, useful for t
 PT ovarian cancer, and as probes, primers, and markers of cancer
 PT progression.
 XX Example 1; Page 253; 378pp; English.
 PS The invention comprises compositions used for the therapy and dia
 CC of ovarian cancer. The compositions comprise one or more ovarian
 CC proteins, their associated polynucleotides, or immunogenic portic
 CC the proteins. The ovarian tumour polynucleotides and polypeptides
 CC useful for stimulating and/or expanding T cells specific for a tu
 CC protein. They are also useful for inhibiting the development of c
 CC a patient with an ovarian tumour DNA or protein by incubating asc
 CC cells allowing them to proliferate, and administering to the pati
 CC sequences can be used as markers for cancer, for example, to moni
 CC ovarian cancer progression. Probes and primers are useful in nucl
 CC hybridisation, in detecting the presence of complementary sequen
 CC given sample, for preparing mutant species and for preparing othe
 CC genetic constructions. Sequences AAG23820-AAS25231 and AAS2528-
 CC represent human ovarian tumour protein cDNA clones
 XX Sequence 281 BP; 98 A; 49 C; 59 G; 74 T; 0 U; 1 Other;
 SQ Query Match 1.9%; Score 26; DB 4; Length 281;
 Best Local Similarity 100.0%; Pred. No. 0.11;
 Matches 26; Conservative 0; Mismatches 0; Indels 0;
 QY 1 ATGTCATTGTTAGACTTTGAAATTC 26
 |||||
 DB 220 ATGTCATTGTTAGACTTTGAAATTC 245
 RESULT 40
 ABQ60530
 ID ABQ60530 standard; cDNA; 386 BP.
 XX AC ABQ60530;
 XX DT 02-AUG-2002 (first entry)
 XX Human colon cancer related nucleotide sequence SEQ ID NO:4225.
 DE Human; colon cancer; cancer; tissue profiling; forensic; mapping
 KW genetic analysis; diagnostic; antisense therapy; gene; ss.
 XX Homo sapiens.
 OS WO200229086-A2.
 PN

001WO-US030732.
 000US-0237271P.
 CORP.
 ste JH, Carroll E, Catino TJ, Dwivedi P, Molino GA,
 Lewis ME;
 15/45.
 nucleic acid that is differentially expressed in cancer
 for determining the presence of colon cancer in a cell or
 and in antisense therapy.
 796pp; English.
 3050787 represent isolated nucleic acids (I) differentially
 cancer tissues. AB878993 to ABB79004 represents proteins
 ABQ60776 to ABQ60787 nucleic acid sequences. (I) can be
 use therapy. An antibody immunoreactive with a polypeptide
 is useful for detecting cancer in a patient sample, and
 the presence or absence of a polynucleotide encoded by a
 which hybridises to (I) in a cell. A probe/primer derived
 be used for determining the presence of a nucleic acid which
 (I), and for determining the phenotype of cells in a sample
 a patient. (I) is useful for determining the presence of
 in a cell or tissue type, for determining the presence of
 type of cancer, in antisense therapy, to generate
 a solid surface, to identify a chromosome on which the
 gene resides, and in tissue profiling, forensics, genetic
 ing and diagnostic applications. (I) can be used to raise
 id to screen for peptide analogues and antagonists
 3P; 128 A; 72 C; 85 G; 99 T; 0 U; 2 Other;
 1.9%; Score 26; DB 6; Length 386;
 100.0%; Pred. No. 0.11;
 0; Mismatches 0; Indels 0; Gaps 0;
 2ATTGTTAGACTTTGAAATTC 26
 2ATTGTTAGACTTTGAAATTC 343
 iard; cDNA; 391 BP.
 (first entry)
 tumour associated polynucleotide sequence SEQ ID NO:961.
 a tumour; ovarian cancer; diagnosis; gene therapy;
 vaccine; ss.
 2.
 2001WO-US001575.
 2000US-0176722P.
 A CORP.

PI Algate PA;
 XX WPI; 2001-425866/45.
 XX Novel ovarian tumor proteins, and nucleic acids encoding them, usi
 PT treat and diagnose cancers, particularly ovarian cancer.
 XX
 PS Claim 5; Page 239; 338pp; English.
 XX
 CC AAH82377 to AAH83678 represent human ovarian tumour-associated
 CC polynucleotide sequences which encode ovarian tumour proteins. The
 CC ovarian tumour protein and polynucleotide sequences have cytostat
 CC activity, and can be used in gene therapy and vaccine production.
 CC ovarian tumour proteins and polynucleotides can be used to inhibi
 CC development of cancer, particularly ovarian cancer. They can also
 CC to diagnose the onset and progression of cancer
 XX Sequence 391 BP; 118 A; 80 C; 88 G; 97 T; 0 U; 8 Other;
 SQ
 Query Match 1.9%; Score 26; DB 5; Length 391;
 Best Local Similarity 100.0%; Pred. No. 0.11;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; G
 QY 1 ATGTCATTGTTAGACTTTGAAATTC 26
 Db 220 ATGTCATTGTTAGACTTTGAAATTC 245
 RESULT 42
 ABX74646/c
 ID ABX74646 standard; cDNA; 425 BP.
 XX
 AC ABX74646;
 XX
 DT 21-MAR-2003 (first entry)
 XX
 DE Human cDNA sequence #113 up-regulated in CC-RCC patients.
 XX
 KW Human; microarray; solid surface; immobilised probe; CC-RCC;
 KW differential expression profile; aggressive CC-RCC tumour type;
 KW non-aggressive CC-RCC tumour type; clear cell renal carcinoma;
 KW gene expression profiling; tumour tissue; gene; ss.
 XX
 OS Homo sapiens.
 XX
 PX WO200279411-A2.
 XX
 PD 10-OCT-2002.
 XX
 XX 29-MAR-2002; 2002WO-US009576.
 XX
 XX 29-MAR-2001; 2001US-0279411P.
 XX
 XX (VAND-) VAN ANDEL INST.
 XX
 XX Haab B, Rhodes D, Teh ET, Takashi M;
 PI WPI; 2003-040679/03.
 DR
 XX New microarray, comprising a matrix of cDNA probe from a set of p
 PT immobilized to a solid surface in predetermined order, useful in
 PT prognosis of patients with clear cell renal carcinoma.
 XX
 XX Claim 35; SEQ ID NO 223; 179pp; English.
 PS
 CC The present invention relates to a microarray comprising a matrix
 CC least one cDNA probe from a set of probes immobilised to a solid
 CC in a predetermined order, where a row of pixels corresponds to re
 CC of one distinct probe from the set. The probes are complementary
 CC nucleic acid sequences that are expressed differentially in aggre
 CC compared to non-aggressive types of clear cell renal carcinoma (C
 CC and that hybridise to the probes under high stringency conditions
 CC microarray is useful for the prognosis of patients with CC-RCC, w

06:25:14 2004

us-09-245-198a-3.oligo.rng

id non-aggressive CC-RCC tumour types are characterised by expression profiles of genes that hybridise with one or more listed on the microarray. The arrays are useful for gene profiling of tumour and normal tissues. The present sequence human cDNA sequence up-regulated in CC-RCC patients

BP; 118 A; 91 C; 78 G; 138 T; 0 U; 0 Other;

arity 1.9%; Score 26; DB 7; Length 425;

conservative 0; Mismatches 0; Indels 0; Gaps 0;

TCATTGTTAGACTTGAATTC 26

TCATTGTTAGACTTGAATTC 45

idard; DNA; 439 BP.

(first entry)

arcinoma related gene sequence SEQ ID NO:1027.

colon; breast; ovary; oesophagus; kidney; thyroid; prostate; pancreas; carcinoma; antitumour; cancerous; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;

A2.

2001WO-US010838.

2000US-0209473P.

2000US-0209531P.

2000US-0233133P.

2000US-0233617P.

2000US-0234009P.

2000US-0234034P.

2000US-0234052P.

2000US-0234509P.

2000US-0234567P.

2000US-0234923P.

2000US-0234924P.

2000US-0235077P.

2000US-0235082P.

2000US-0235134P.

2000US-0235280P.

2000US-0235637P.

2000US-0235638P.

2000US-0235711P.

2000US-0235720P.

2000US-0235840P.

2000US-0235863P.

2000US-0236028P.

2000US-0236032P.

2000US-0236033P.

2000US-0236034P.

2000US-0236109P.

2000US-0236111P.

2000US-0236842P.

2000US-0236891P.

2000US-0237172P.

2000US-0237173P.

2000US-0237278P.

2000US-0237294P.

02-OCT-2000; 2000US-0237295P.

02-OCT-2000; 2000US-0237316P.

03-OCT-2000; 2000US-0237425P.

03-OCT-2000; 2000US-0237598P.

03-OCT-2000; 2000US-0237604P.

03-OCT-2000; 2000US-0237608P.

03-OCT-2000; 2000US-0237608P.

01-NOV-2000; 2000US-0244867P.

01-NOV-2000; 2000US-0245084P.

(AVAL-) AVALON PHARM.

Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrig-

Soppet DR, Weaver Z;

WPI; 2002-188264/24.

Screening for anti-neoplastic agent involves exposing cells to a agent to be tested for anti-neoplastic activity, and determining in expression of a gene of a signature gene set.

Claim 1; SEQ ID NO 1027; 44pp; English.

The present invention describes a method (M1) for screening for neoplastic agent. The method involves exposing cells to a chemi- to be tested for anti-neoplastic activity, determining a change expression of at least one gene (I) of a signature gene set, whe comprises a sequence (S) selected from 8447 sequences (given in to ABL70110), or is at least 95% identical to (S), where a chang- expression is indicative of anti-neoplastic activity. (I) has cy activity and can be used in gene therapy. M1 can be used for scr anti-neoplastic agent, and can be used for producing a product w the data collected with respect to the anti-neoplastic agent as of M1, and the data is sufficient to convey the chemical structu properties of the agent. M1 can be used in the treatment of canc as colon, breast, stomach, lung, thyroid, oesophageal, ovarian, prostate or pancreatic cancer, adenocarcinoma, carcinoma, clear cancer, infiltrating ductal cancer, infiltrating lobular cancer, cell carcinoma, neuroendocrine carcinoma, papillary carcinoma an tumour

Sequence 439 BP; 118 A; 101 C; 83 G; 137 T; 0 U; 0 Other;

Query Match 1.9%; Score 26; DB 6; Length 439;

Best Local Similarity 100.0%; Pred. No. 0.11;

Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 1 ATGTCATTGTTAGACTTGAATTC 26

DB 72 ATGTCATTGTTAGACTTGAATTC 47

RESULT 44

ABN96246/C

ID ABN96246 standard; DNA; 439 BP.

XX AC ABN96246;

XX XX

DT 13-AUG-2002 (first entry)

XX Gene #2744 used to diagnose liver cancer.

XX Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic; metastatic liver tumour; cytostatic; expression profile; disease disease progression; drug toxicity; drug efficacy; drug metaboli

OS Homo sapiens.

XX WO200229103-A2.

XX PD 11-APR-2002.

XX 02-OCT-2001; 2001WO-US030589.

PF

2000US-0237054P.
LOGIC INC.
ares C, Peres-Da-Silva S, Vockley JG;
119/45.
a detecting the progression of liver cancer, hepatocellular
metastatic liver tumor in a patient, involves detecting the
ession of two or more genes in a liver tissue sample.
ID NO 2744; 298pp; English.
relates to a novel method for diagnosing and detecting the
f liver cancer, hepatocellular carcinoma or metastatic liver
atient, and differentiating metastatic liver cancer from
r carcinoma in a patient, involving detecting the level of
two or more genes represented in ABN93503-ABN97455 in a
The method of the invention has hepatotropic, and
ivity. The method is useful for diagnosing and detecting
on of liver cancer, hepatocellular carcinoma and metastatic
na in a patient. The method is useful for identifying
files which serve as useful diagnostic markers as well as
can be used to monitor disease states, disease progression,
drug efficacy and drug metabolism. Note: The sequence data
at did not form part of the printed specification, but was
electronic format directly from WIPO at
pub/published_pct_sequences
BP; 118 A; 101 C; 83 G; 137 T; 0 U; 0 Other;
1.9%; Score 26; DB 6; Length 439;
arity 100.0%; Pred. No. 0.11;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
CATGTAGACTTTGAAATTC 26
CATGTAGACTTTGAAATTC 47
iard; cDNA; 452 BP.
(first entry)
a protein 5' EST, SEQ ID NO: 10997.
; expressed sequence tag; secreted protein; cDNA isolation;
chromosome mapping; ss.
2000EP-00200610.
99US-0122487P.
F.
iwards J, Duclert A, Giordano J;
381/45.
id that is a 5' expressed sequence tag (5' EST) for
as and genomic DNAs that correspond to 5'ESTs and for

PT diagnostic, forensic, gene therapy and chromosome mapping procedu
XX Claim 1; SEQ ID NO 10997; 71pp + Sequence Listing; English.
PS
XX
CC The present sequence is one of a large number of 5' ESTs derived
CC mRNAs encoding secreted proteins. NO ORF has yet been conclusivel
CC identified within the present sequence. The 5' ESTs were prepared
CC total human RNAs or polyA+ RNAs derived from 30 different tissues
CC sequences usually correspond mainly to the 3' untranslated region
CC of the mRNA because they are often obtained from oligo-dT primed
CC libraries. Such ESTs are not well suited for isolating cDNA sequ
CC derived from the 5' ends of mRNAs and even in those cases where l
CC cDNA sequences have been obtained, the full 5' UTR is rarely incl
CC ESTs are derived from mRNAs with intact 5' ends and can therefore
CC to obtain full length cDNAs and genomic DNAs. 5' ESTs are also us
CC diagnostic, forensic, gene therapy and chromosome mapping procedu
CC They are used to obtain upstream regulatory sequences and to desi
CC expression and secretion vectors
XX
SQ Sequence 452 BP; 122 A; 95 C; 112 G; 122 T; 0 U; 1 Other;
Query Match 1.9%; Score 26; DB 3; Length 452;
Best Local Similarity 100.0%; Pred. No. 0.11;
Matches 26; Conservative 0; Mismatches 0; Indels 0; G
QY 1 ATGTCATTGTTAGACTTTGAAATTC 26
Db 414 ATGTCATTGTTAGACTTTGAAATTC 439
RESULT 46
ABV86720
ID ABV86720 standard; cDNA; 469 BP.
XX
AC ABV86720;
XX
DT 13-DEC-2002 (first entry)
XX Human colon cancer related cDNA SEQ ID NO 31.
XX Human; colon; cancer; cytostatic; tumour; gene therapy; vaccine;
KW ss.
XX Homo sapiens.
OS
XX WO200258534-A2.
PN
XX 01-AUG-2002.
XX
XX 16-NOV-2001; 2001WO-US043704.
PF
XX 20-NOV-2000; 2000US-0252222P.
PR
XX 06-FEB-2001; 2001US-0267011P.
PR
XX 28-MAR-2001; 2001US-0279670P.
PR
XX 10-JUL-2001; 2001US-0304037P.
XX
PA (CORI-) CORIXA CORP.
XX
XX Stolk JA, Xu J, Chenault RA, Meagher MJ, Secrist H, King GE;
PI
XX WPI; 2002-608400/65.
DR
XX New isolated tumor colon polynucleotide and polypeptide, useful f
PT diagnosis, prevention and/or treatment of cancer, in particular c
PT cancer.
XX
XX Claim 1; SEQ ID NO 31; 266pp + Sequence Listing; English.
PS
XX The invention relates to a human colon tumour expressed polynucle
CC (i) encoding a polypeptide (ii, ABP67991-ABP67996) comprising: (i
CC 2600 fully defined nucleotide sequences (ABV8669-ABV89289); (ii)
CC complements of (i); (iii) at least 20 contiguous residues of (i);
CC sequences that hybridize to (i), under moderately stringent condi

s having at least 75% or 90% identity to (i); or (vi) variants of (i). The compositions and methods of the present invention are useful for the diagnosis, prevention and/or treatment of colon cancer. (i) can be used in gene therapy and are useful in pharmaceutical compositions such as vaccines. Sequence data for this patent did not form part of the printed matter, but was obtained in electronic format directly from WIPO int/pub/published_pct_sequences

BP; 142 A; 94 C; 111 G; 122 T; 0 U; 0 Other;

1.9%; Score 26; DB 6; Length 469;

larity 100.0%; Pred. No. 0.11; 0; Indels 0; Gaps 0; Conservative 0; Mismatches 0;

TCATTGTTAGACTTTGAAATTC 26

TCATTGTTAGACTTTGAAATTC 425

ndard; cDNA; 483 BP.

(first entry)

cancer-associated cDNA, SEQ ID No 558.

cancer; immunogenic; vaccine; tumour; gene; ss.

A2.

2001WO-US023826.

2000US-0223265P.

2000US-0237406P.

2001US-0277495P.

2001US-0302702P.

XA CORP.

J, Secret H;

7462/30.

cleotide encoding colon tumor polypeptides, useful as treating colon cancers.

309; 425pp; English.

a relates to isolated polynucleotides (I) encoding colon peptides (II). (I) is useful for stimulating an immune patient and treating colon cancer in a patient. Ideas derived from (I) are useful for determining the presence a patient. (I) and (II) are useful in pharmaceutical, e.g. vaccines, and other compositions for the diagnosis and colon cancer. A composition comprising a first component physiologically acceptable carriers and immunostimulants, en-presenting cell expressing (II) is useful for inhibiting of cancer in a patient. (I) is useful in the design and of ribozyme molecules for inhibiting expression of tumour and (I). ABK54531-ABK5464 represent human colon cancer cDNA the invention

BP; 155 A; 97 C; 112 G; 119 T; 0 U; 0 Other;

1.9%; Score 26; DB 6; Length 483;

Best Local Similarity 100.0%; Pred. No. 0.11; Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 1 ATGTCATTGTTAGACTTTGAAATTC 26

Db 397 ATGTCATTGTTAGACTTTGAAATTC 422

RESULT 48

AAZ51563

ID AAZ51563 standard; cDNA; 486 BP.

XX AAZ51563;

AC AAZ51563;

XX 21-JUN-2000 (first entry)

DT Human hypoxia response regulating gene, 77H4 related cDNA clone

XX Hypoxia response regulating gene; gene 77H4; human; EST 3D; angi cardiant; apoptosis; vasotropic; cytostatic; ophthalmological; s cerebroprotective; antagonist; regulator; inhibitor; treatment; KW hypoxia associated pathology; HAP; gene therapy; diagnosis; reti steroid receptor coactivator; SRA; ischaemia; myocardial infarct XX Homo sapiens.

XX Key Location/Qualifiers

FT polyA_signal 449..454

FT /*tag= a

XX WO200012525-A1.

PN 09-MAR-2000.

XX 27-AUG-1999; 99WO-US020394.

XX 27-AUG-1998; 98US-0098158P.

PR 05-MAY-1999; 99US-0132684P.

XX (QUAR-) QUARK BIOTECH INC.

PA (KOHN/) KOHN K.

XX Einat P, Skalter R, Feinstein E;

XX WPI; 2000-256577/22.

XX Novel polynucleotides capable of regulating angiogenesis or apop useful for diagnosis and treatment of hypoxia, ischemia and tumo

XX Claim 1; Fig 7b; 78pp; English.

XX The present sequence is the human hypoxia response regulating ge related cDNA clone 3D. The gene 77H4 has similarity to steroid r transcriptional co-activator, SRA function and can serve as a co in some transcriptional complexes. It has vasotropic, cardiant, CC ophthalmological, cytostatic and cerebroprotective activity. Ant CC of the encoded protein, functions as a regulator of apoptosis or CC angiogenesis. The protein encoded by this polynucleotide, the CC biologically active product from enzymatic activity of the prote CC inhibitor of the enzymatic activity is useful for regulating hyp CC associated pathologies (HAP). It is useful for gene therapy, dia CC and treatment of tumour growth and ischaemia, e.g., retinopathy, CC myocardial infarction and stroke

XX SQ Sequence 486 BP; 160 A; 92 C; 113 G; 121 T; 0 U; 0 Other;

Query Match 1.9%; Score 26; DB 3; Length 486;

Best Local Similarity 100.0%; Pred. No. 0.11;

Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 1 ATGTCATTGTTAGACTTTGAAATTC 26

Db 400 ATGTCATTGTTAGACTTTGAAATTC 425

DE Human endothelial cell cDNA #4820.
XX Human; ss; sequencing by hybridisation; SBH; expressed sequence t
KW genome mapping; biodiversity; genetic disorder.
XX Homo sapiens.
XX US2003073623-A1.
XX 17-APR-2003.
XX 30-JUL-2001; 2001US-00918995.
XX 30-JUL-2001; 2001US-00918995.
XX (DRMA/) DRMANAC R T.
XX (LABAT/) LABAT I.
XX (STAC/) STACHE-CRAIN B.
XX (DICK/) DICKSON M C.
XX (JONE/) JONES L W.
XX Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;
XX WPI; 2003-615964/58.
XX New polynucleotide sequences obtained from various cDNA libraries
XX as hybridization probes, as oligomers for PCR, for chromosome and
XX mapping, in the recombinant production of protein, or in generati
XX antisense DNA or RNA.
XX Claim 1; SEQ ID NO 23899; 44pp; English.
XX The invention relates to an isolated polynucleotide comprising an
XX 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose seque
XX determined by the technique of SBH (sequencing by hybridisation).
XX included is a purified polypeptide comprising a sequence correspo
XX a reading frame of the novel polynucleotide. The nucleic acid seq
XX are useful in diagnostics as expressed sequence tags (EST) for
XX identifying expressed genes or for physical mapping of the human
XX in forensics, in assessing biodiversity, or in identifying muta
XX responsible for genetic disorders and other traits. The nucleotid
XX sequences are also useful as hybridisation probes, as oligomers f
XX for chromosome and gene mapping, in the recombinant production of
XX protein, or in generating antisense DNA or RNA. The purified poly
XX is useful for generating antibodies specific for it. The present
XX is one of the 38043 isolated cDNA/EST sequences. Note: the sequen
XX for this patent did not form part of the printed specification, b
XX obtained in electronic format directly from USPTO at
XX seqdata.uspto.gov/sequence.html?DocID=20030073623
XX Sequence 498 BP; 152 A; 119 C; 117 G; 107 T; 0 U; 3 Other;
SQ Query Match 1.9%; Score 26; DB 8; Length 498;
Best Local Similarity 100.0%; Pred. No. 0.11;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps
QY 1 AGTCTATTCTTACACTTTGAAATTTTC 26
Db 349 AGTCTATTCTTACACTTTGAAATTTTC 374
RESULT 51
AAC01272
ID AAC01272 standard; cDNA; 516 BP.
XX AAC01272;
AC AAC01272;
DT 06-OCT-2000 (first entry)
XX Human secreted protein 5' EST, SEQ ID NO: 1270.
XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA iso
KW gene therapy; chromosome mapping; ss.

DE Human endothelial cell cDNA #4820.
XX Human; ss; sequencing by hybridisation; SBH; expressed sequence t
KW genome mapping; biodiversity; genetic disorder.
XX Homo sapiens.
XX US2003073623-A1.
XX 17-APR-2003.
XX 30-JUL-2001; 2001US-00918995.
XX 30-JUL-2001; 2001US-00918995.
XX (DRMA/) DRMANAC R T.
XX (LABAT/) LABAT I.
XX (STAC/) STACHE-CRAIN B.
XX (DICK/) DICKSON M C.
XX (JONE/) JONES L W.
XX Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;
XX WPI; 2003-615964/58.
XX New polynucleotide sequences obtained from various cDNA libraries
XX as hybridization probes, as oligomers for PCR, for chromosome and
XX mapping, in the recombinant production of protein, or in generati
XX antisense DNA or RNA.
XX Claim 1; SEQ ID NO 23899; 44pp; English.
XX The invention relates to an isolated polynucleotide comprising an
XX 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose seque
XX determined by the technique of SBH (sequencing by hybridisation).
XX included is a purified polypeptide comprising a sequence correspo
XX a reading frame of the novel polynucleotide. The nucleic acid seq
XX are useful in diagnostics as expressed sequence tags (EST) for
XX identifying expressed genes or for physical mapping of the human
XX in forensics, in assessing biodiversity, or in identifying muta
XX responsible for genetic disorders and other traits. The nucleotid
XX sequences are also useful as hybridisation probes, as oligomers f
XX for chromosome and gene mapping, in the recombinant production of
XX protein, or in generating antisense DNA or RNA. The purified poly
XX is useful for generating antibodies specific for it. The present
XX is one of the 38043 isolated cDNA/EST sequences. Note: the sequen
XX for this patent did not form part of the printed specification, b
XX obtained in electronic format directly from USPTO at
XX seqdata.uspto.gov/sequence.html?DocID=20030073623
XX Sequence 498 BP; 152 A; 119 C; 117 G; 107 T; 0 U; 3 Other;
SQ Query Match 1.9%; Score 26; DB 8; Length 498;
Best Local Similarity 100.0%; Pred. No. 0.11;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps
QY 1 AGTCTATTCTTACACTTTGAAATTTTC 26
Db 349 AGTCTATTCTTACACTTTGAAATTTTC 374
RESULT 51
AAC01272
ID AAC01272 standard; cDNA; 516 BP.
XX AAC01272;
AC AAC01272;
DT 06-OCT-2000 (first entry)
XX Human secreted protein 5' EST, SEQ ID NO: 1270.
XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA iso
KW gene therapy; chromosome mapping; ss.

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PR 03-JUL-2001; 2001US-0302702P.
XX
XX {CORI-) CORIXA CORP.
XX
XX Pyle RA, Xu J, Secrist H;
XX
XX WPI; 2002-257462/30.
XX
XX Novel polynucleotide encoding colon tumor polypeptides, useful as
XX PT vaccines for treating colon cancers.
XX PT
XX Claim 1; Page 403; 425pp; English.
XX PS
XX
XX The invention relates to isolated polynucleotides (I) encoding cc
XX CC tumour polypeptides (II). (I) is useful for stimulating an immune
XX CC response in a patient and treating colon cancer in a patient.
XX CC Oligonucleotides derived from (I) and (II) are useful for determining the
XX CC of cancer in a patient. (I) and (II) are useful in pharmaceutical
XX CC compositions, e.g. vaccines, and other compositions for the diag-
XX CC nosis and treatment of colon cancer. A composition comprising a first compo-
XX CC nent selected from physiologically acceptable carriers and immunostimu-
XX CC lants and an antigen-presenting cell expressing (II) is useful for inhi-
XX CC bition of development of cancer in a patient. (I) is useful in the design of
XX CC a preparation of ribozyme molecules for inhibiting expression of tu-
XX CC mor polypeptides and (I). ABK54531-ABK5464 represent human colon car-
XX CC cinoma sequences of the invention
XX
XX Sequence 531 BP; 153 A; 105 C; 128 G; 143 T; 0 U; 2 Other;
XX SQ
Query Match 1.9%; Score 26; DB 6; Length 531;
Best Local Similarity 100.0%; Pred. No. 0.11;
Matches 26; Conservative 0; Mismatches 0; Indels 0; G
QY 1 ATGTCATTGTTAGACTTTGAAATTTC 26
DB |||||||||||||||||||||||||
459 ATGTCATTGTTAGACTTTGAAATTTC 484
RESULT 53
AAC01271
ID AAC01271 standard; cDNA; 540 BP.
XX
XX AAC01271;
AC
AC
DT 06-OCT-2000 (first entry)
DE Human secreted protein 5' EST, SEQ ID NO: 1269.
XX
XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA iso-
XX KW gene therapy; chromosome mapping; ss.
XX KW Homo sapiens.
XX OS
XX EP1033401-A2.
XX PN
XX 06-SEP-2000.
XX
XX 21-FEB-2000; 2000EP-00200610.
XX PF
XX 26-FEB-1999; 99US-0122487P.
XX PR
XX (GEST ) GENSET.
XX
XX Dumas Milne Edwards J, Duclert A, Giordano J;
XX PA
XX WPI; 2000-500381/45.
XX DR
XX P-PSDB; AAG01265.
XX
XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
XX PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and fo-
XX PT rmal diagnostic, forensic, gene therapy and chromosome mapping procedu-
XX PT res.
XX PS Claim 1; SEQ ID NO 1269; 71pp + Sequence Listing; English.

```

2000EP-00200610.
99US-0122487P.
T.
dwards J, Duclert A, Giordano J;
381/45.
266.
acid that is a 5' expressed sequence tag (5' EST) for
cDNAs and genomic DNAs that correspond to 5' ESTs and for
forensic, gene therapy and chromosome mapping procedures.
ID NO 1270; 71pp + Sequence Listing; English.
sequence is one of a large number of 5' ESTs derived from
secreted proteins. An ORF has been identified within the
5' ESTs were prepared from total human RNAs or polyA+ RNAs
30 different tissues. EST sequences usually correspond
3' untranslated region (UTR) of the mRNA because they are
d from oligo-dT primed cDNA libraries. Such ESTs are not
or isolating cDNA sequences derived from the 5' ends of
n in those cases where longer cDNA sequences have been
full 5' UTR is rarely included. 5' ESTs are derived from
tact 5' ends and can therefore be used to obtain full length
omic DNAs. 5' ESTs are also used in diagnostic, forensic,
and chromosome mapping procedures. They are used to obtain
latory sequences and to design expression and secretion
BP; 134 A; 106 C; 131 G; 145 T; 0 U; 0 Other;
1.9%; Score 26; DB 3; Length 516;
arity 100.0%; Pred.No. 0.11;
conservative 0; Mismatches 0; Indels 0; Gaps 0;
CATGTTAGACTTTGAAATTC 26
|||||
CATGTTAGACTTTGAAATTC 503
dard; cDNA; 531 BP.
(first entry)
ancer-associated cDNA, SEQ ID No 877.
cancer; immunogenic; vaccine; tumour; gene; ss.
2.
2001WO-US023826.
2000US-0223265P.
2000US-0237406P.
2001US-0277495P.

sequence is one of a large number of 5' ESTs derived from 3 secreted proteins. An ORF has been identified within the 5' ESTs were prepared from total human RNAs or polyA+ RNAs 30 different tissues. EST sequences usually correspond to 3' untranslated region (UTR) of the mRNA because they are derived from oligo-dT primed cDNA libraries. Such ESTs are not or isolating cDNA sequences derived from the 5' ends of a in those cases where longer cDNA sequences have been full 5' UTR is rarely included. 5' ESTs are derived from tect 5' ends and can therefore be used to obtain full length cmic DNAs. 5' ESTs are also used in diagnostic, forensic, and chromosome mapping procedures. They are used to obtain latory sequences and to design expression and secretion

3P; 135 A; 110 C; 137 G; 152 T; 0 U; 6 Other;

arity 1.9%; Score 26; DB 3; Length 540;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

TATGTTAGACTTTGAAATTC 26

TATGTTAGACTTTGAAATTC 527

lard; cDNA; 570 BP.

first entry)

associated gene sequence SEQ ID NO:292.

associated gene; cancer antigen; detection; cancer; ostatic; proliferative; vulnary; immunomodulator; antisthmatic; antirheumatic; antiarthritic; antiviral; ry; antithyroid; antiallergic; antibacterial; cardiant; ; neuroprotective; thrombolytic; coagulant; neutropic; tipsporadic; antiangiogenic; gene therapy; inflammation; r; haematopoietic cell disorder; autoimmune disorder; ion; graft versus host disease; organ rejection; hrombolytic; cardiovascular disorder; infection; isease; drug screening; ss.

000WO-US005882.

99US-0124270P.

GENOME SCI INC.

en SM;

33/55.

89.

nucleic acids comprising sequences encoding peptides ating or diagnosing e.g. cancer.

853; 2352pp; English.

C78448 encode the human cancer associated proteins given in B44239. The proteins can have activities based on the lls the genes are expressed in. Example of activities

CC include: cytostatic; proliferative; vulnary; immunomodulator; CC anti-diabetic; antiasthmatic; antirheumatic; antiarthritic; CC anti-inflammatory; antithyroid; antiallergic; antibacterial; anti dermatological; neuroprotective; cardiant; thrombolytic; coagular CC neutropic; vasotropic; antiporatic and antiangiogenic. The CC polynucleotides and polypeptides can be used for preventing, trea CC ameliorating medical conditions and diagnosing pathological condi CC Polynucleotides, polypeptides, antibodies, agonists and antagonis CC the present invention may be used to treat immune disorders by ac CC or inhibiting the proliferation, differentiation or mobilisation CC immune cells, to treat disorders of haematopoietic cells, autoimm CC disorders, allergic reactions, graft versus host disease and orga CC rejection, modulate haemostatic or thrombolytic activity, modulat CC inflammation, cancers, cardiovascular disorders, neurological dis CC bacterial or viral infections. The peptides, nucleotides, antibod CC agonists and antagonists may be also be used in drug screens. AAC CC AAC78457 and AAB44240 represent sequences used in the exemplifica CC the present invention XX

SQ Sequence 570 BP; 171 A; 112 C; 139 G; 145 T; 0 U; 3 Other;

Query Match 1.9%; Score 26; DB 3; Length 570;

Best Local Similarity 100.0%; Pred. No. 0.11;

Matches 26; Conservative 0; Mismatches 0; Indels 0; G

QY 1 ATGTCATGTTAGACTTTGAAATTC 26

Db 469 ATGTCATGTTAGACTTTGAAATTC 494

RESULT 55

AAZ51562

ID AAZ51562 standard; cDNA; 580 BP.

XX AAZ51562;

XX 21-JUN-2000 (first entry)

Human hypoxia response regulating gene, 77H4 related cDNA clone 1;

XX Hypoxia response regulating gene; gene 77H4; human; EST 18E; card. KW apoptosis; angiogenesis; vasotropic; cytostatic; ophthalmological KW cerebroprotective; antagonist; regulator; inhibitor; treatment; ti KW hypoxia associated pathology; HAP; gene therapy; diagnosis; ischa KW steroid receptor coactivator; SRA; retinopathy; myocardial infarci KW stroke; ss.

XX Homo sapiens.

XX Key Location/Qualifiers

FT polyA_signal 536..541

FT /*tag= a

XX WO200012525-A1.

XX 09-MAR-2000.

XX 27-AUG-1999; 99WO-US020394.

XX 27-AUG-1999; 98US-0098158P.

XX 05-MAY-1999; 99US-0132684P.

XX (QUAR-) QUARK BIOTECH INC.

XX (KOHN/) KOHN K.

XX Einat P, Skaliter R, Feinstein E;

XX WPI; 2000-256577/22.

XX Novel polynucleotides capable of regulating angiogenesis or apoptc PT useful for diagnosis and treatment of hypoxia, ischemia and tumor XX Claim 1; Fig 7a; 78pp; English.

sequence is the human hypoxia response regulating gene, 77H4, clone 18F. The gene 77H4 has similarity to steroid receptor nuclear co-activator, SRA function and can serve as a coactivator transcriptional complexes. It has vasotropic, cardiact, local, cytostatic and cerebroprotective activity. Antagonist of protein, functions as a regulator of apoptosis or active product from enzymatic activity of the protein or the enzymatic activity is useful for regulating hypoxia pathologies (HAP). It is useful for gene therapy, diagnosis of tumour growth and ischaemia, e.g., retinopathy, infarction and stroke

BP; 177 A; 111 C; 134 G; 158 T; 0 U; 0 Other;

1.9%; Score 26; DB 3; Length 580;

Best Local Similarity 100.0%; Pred. No. 0.11; 0; Indels 0; Gaps 0; Mismatches 0; Conservative 0;

ATGTCATTGTTAGACTTTGAAATTC 26

|||||
ATGTCATTGTTAGACTTTGAAATTC 512

Standard; cDNA; 580 BP.

(first entry)

744 gene sequence.

ia-regulated activity; neurotoxic stress; hypoxia; ischaemia; ois; angiogenesis; cerebroprotective; gene therapy; inhibitor of oxidative stress-mediated apoptosis; angiogenesis; gene 7744; gene; ss.

Al.

2001US-00802472.

97US-0056453P.

98US-00136109.

98US-0098158P.

99US-0132684P.

99US-00384096.

P.

TER R.

STEIN E.

Alter R, Feinstein E;

474/02.

polypeptides and genes associated with hypoxia-regulated for treating stroke, hypoxia and ischemia.

61; 72bp; English.

vention relates to a new polypeptide associated with ated activity. The invention is useful in diagnostic assays. is further useful as a diagnostic tool which can be used to ir presence in a cell. The invention is also useful for odies that could be used in diagnostic assays for the the protein and for determining if any given cell had been

CC subjected to neurotoxic stress. The invention can be used to pro
CC neural cells from, and ameliorate the effects of, hypoxia and is
CC and thus in the treatment of stroke, hypoxia and ischaemia. The
CC is also useful to prevent apoptosis and promote angiogenesis. Th
CC invention can be used in diagnostic assays for cells that have b
CC subjected to hypoxia or ischaemia, and in screening assays to id
CC agents capable of enhancing gene expression. The present nucleic
CC sequence represents the human gene 7744 gene of the invention. N
CC specification states that this sequence encodes the human gene 7
CC protein (ABG71808) but this does not appear to be the case
XX

SQ Sequence 580 BP; 177 A; 111 C; 134 G; 158 T; 0 U; 0 Other;

Query Match

Best Local Similarity 1.9%; Score 26; DB 7; Length 580;

Matches 26; Conservative 0; Mismatches 0; Indels 0;

QY 1 ATGTCATTGTTAGACTTTGAAATTC 26

|||||
Db 487 ATGTCATTGTTAGACTTTGAAATTC 512

RESULT 57

ABQ58318/c

ID ABQ58318 standard; cDNA; 400 BP.

XX AC ABQ58318;

XX 02-AUG-2002 (first entry)

DE Human colon cancer related nucleotide sequence SEQ ID NO:2013.

XX Human; colon cancer; cancer; tissue profiling; forensic; mapping
KW genetic analysis; diagnostic; antisense therapy; gene; ss.

XX Homo sapiens.

OS WO200229086-A2.

PN 11-APR-2002.

PF 02-OCT-2001; 2001WO-US030732.

PR 02-OCT-2000; 2000US-0237271P.

PA (FARB) BAYER CORP.

PI Burgess C, Astle JH, Carroll E, Catino TJ, Dwivedi P, Molin
PI Thaglingam A, Lewis ME;

WPI; 2002-426115/45.

PT New isolated nucleic acid that is differentially expressed in car
PT tissues useful for determining the presence of colon cancer in a
PT tissue type, and in antisense therapy.

PS Claim 1; Fig 1; 796pp; English.

XX ABQ56306 to ABQ60787 represent isolated nucleic acids (I) differ
CC expressed in cancer tissues. ABQ78993 to ABQ79004 represent prote
CC encoded by the ABQ60776 to ABQ60787 nucleic acid sequences. (I)
CC used in antisense therapy. An antibody immunoreactive with a pol
CC encoded by (I) is useful for detecting cancer in a patient sample
CC for detecting the presence or absence of a polynucleotide encode
CC nucleic acid which hybridises to (I) in a cell. A probe/primer d
CC from (I) can be used for determining the presence of a nucleic ac
CC hybridises to (I), and for determining the phenotype of cells in
CC of cells from a patient. (I) is useful for determining the presen
CC colon cancer in a cell or tissue type, for determining the presen
CC state of other type of cancer, in antisense therapy, to generate
CC macroarrays on a solid surface, to identify a chromosome on which
CC corresponding gene resides, and in tissue profiling, forensics, s
CC analysis, mapping and diagnostic applications. (I) can be used to

06:25:14 2004

us-09-245-198a-3.oligo.rng

nd to screen for peptide analogues and antagonists

BP; 100 A; 80 C; 61 G; 138 T; 0 U; 21 Other;

1.8%; Score 25; DB 6; Length 400;

arity 100.0%; Pred.No. 0.32;

conservative 0; Mismatches 0; Indels 0; Gaps 0;

CAATGTTAGACTTTGAAATTT 25

|||||

CAATGTTAGACTTTGAAATTT 69

dard; cDNA; 626 BP.

(first entry)

novel human diagnostic protein #230.

some mapping; gene mapping; gene therapy; forensic;
nt; medical imaging; diagnostic; genetic disorder; ss.

2.

2001WO-US008631.

2000US-00540217.

2000US-00649167.

INC.

Liu C, Tang YT;

362/73.

239.

polynucleotide and encoded polypeptides, useful in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits and to assess

CD NO 230; 103pp; English.

relates to isolated polynucleotide (I) and polypeptide (II)
is useful as hybridisation probes, polymerase chain
primers, oligomers, and for chromosome and gene mapping,
nant production of (II). The polynucleotides are also used
as expressed sequence tags for identifying expressed
useful in gene therapy techniques to restore normal
(I) or to treat disease states involving (II). (II) is
erating antibodies against it, detecting or quantitating a
tissue, as molecular weight markers and as a food
(I) and its binding partners are useful in medical imaging
using (II). (I) and (II) are useful for treating disorders
rant protein expression or biological activity. The
id polynucleotide sequences have applications in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits to assess biodiversity
other types of data and products dependent on DNA and
ences. AAS4197-AAS94564 represent novel human diagnostic
es of the invention. Note: The sequence data for this
appear in the printed specification, but was obtained in
mat directly from WIPO at
ub/published_pct_sequences

SQ Sequence 626 BP; 173 A; 126 C; 157 G; 170 T; 0 U; 0 Other;

Query Match

1.8%; Score 25; DB 5; Length 626;

Best Local Similarity 100.0%; Pred.No. 0.31;

Matches 25; Conservative 0; Mismatches 0; Indels 0; C

QY 1 ATGTCATTGTTAGACTTTGAAATTT 25

|||||

542 ATGTCATTGTTAGACTTTGAAATTT 566

RESULT 59

AAX23451/c

ID AAX23451 standard; DNA; 24 BP.

XX AC AAX23451;

XX 18-JUN-1999 (first entry)

XX Human TNRL3 RACE primer 2.

XX Tumour necrosis factor receptor; signal transducer molecule; TNF;
XX developmental abnormality; gestational abnormality; prostate ca
XX APO6; APO8; APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy;
XX cytoplasmic domain; immunogen; antibody preparation; breast carci
XX apoptosis; human; primer; ss.

OS Synthetic.

OS Homo sapiens.

XX WO9911791-A2.

XX 11-MAR-1999.

XX 04-SEP-1998; 98WO-US018393.

XX 05-SEP-1997; 97US-00924634.

XX (UNIW) UNIV WASHINGTON.

XX Chaudhary PM;

XX WPI; 1999-20519-1/17.

XX New Tumor Necrosis Factor family receptor polypeptides and ligand
XX useful for diagnosis and treatment of prostate cancer and develop
XX or gestational abnormalities.

XX Example VII; Page 121; 156pp; English.

XX This invention describes isolated Tumor Necrosis Factor (TNF) fam
XX receptor polypeptides: APO4, APO6, APO8 and APO9 or their active
XX fragments, and isolated TNF related ligands 1 and 3 (TNRL1 and TN
XX their active fragments. APO4 is useful for diagnosing prostate ca
XX determining levels of APO4 in an individual. Prostate cancer can
XX treated using APO4 selective binding agents linked to a therapeutic
XX moiety. APO4 polypeptides are also useful for identifying selecti
XX binding agents, useful in diagnosis/treatment of disease by bindi
XX agents to the polypeptide/active fragment which is extracellular,
XX expressed on the cell surface. The binding is preferably performe
XX vivo. APO4 polypeptides/ active fragments are also useful for scr
XX for agonists and antagonists by binding and observing the changer
XX activity. Effective pharmacological agents useful in diagnosis or
XX treatment of disease are also identified using APO4 polypeptides/
XX fragments and APO4 signal transducer molecules that specifically
XX with a cytoplasmic domain of APO4 and detecting a change in level
XX activity. The method is performed in vivo or in vitro. APO polype
XX are all useful as immunogens for preparing antibodies. APO4 is al
XX useful for diagnosis/treatment of developmental or gestational
XX abnormalities. APO8 was transfected to human breast carcinoma cell
XX MCF-7, and induced apoptosis

XX Sequence 24 BP; 7 A; 9 C; 3 G; 5 T; 0 U; 0 Other;

1.7%; Score 24; DB 2; Length 24;
 larity 100.0%; Pred.No.1.1;
 Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GATGAGGGGAAGGCTGCTAC 633
 |||||
 GATGAGGGGAAGGCTGCTAC 1

ndard; DNA; 24 BP.

(first entry)

RACE primer 1.

sis factor receptor; signal transducer molecule; TNF; APO4;
 1 abnormality; Gestational abnormality; prostate cancer;
 APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy; disease;
 domain; immunogen; antibody preparation; breast carcinoma;
 uman; primer; SS.

98WO-US018393.

97US-00924634.

WASHINGTON.

5191/17.

rosis Factor family receptor polypeptides and ligands -
 diagnosis and treatment of prostate cancer and developmental
 abnormalities.

Page 121; 156pp; English.

on describes isolated Tumor Necrosis Factor (TNF) family
 peptides: APO4, APO6, APO8 and APO9 or their active
 and isolated TNF related ligands 1 and 3 (TNRL1 and TNRL3) or
 fragments. APO4 is useful for diagnosing prostate cancer by
 levels of APO4 in an individual. Prostate cancer can also be
 APO4 selective binding agents linked to a therapeutic
 polypeptides are also useful for identifying selective
 is, useful in diagnosis/treatment of disease by binding of
 a polypeptide/active fragment which is extracellular, or
 the cell surface. The binding is preferably performed in
 polypeptides/ active fragments are also useful for screening
 and antagonists by binding and observing the change in APO4
 active pharmacological agents useful in diagnosis or
 disease are also identified using APO4 polypeptides/active
 APO4 signal transducer molecules that specifically interact
 laemic domain of APO4 and detecting a change in level of APO4
 a method is performed in vivo or in vitro. APO polypeptides
 all as immunogens for preparing antibodies. APO4 is also
 diagnosis/treatment of developmental or gestational
 APO8 was transfected to human breast carcinoma cell line
 induced apoptosis

3P; 7 A; 2 C; 13 G; 2 T; 0 U; 0 Other;

Query Match 1.7%; Score 24; DB 2; Length 24;
 Best Local Similarity 100.0%; Pred.No.1.1;
 Matches 24; Conservative 0; Mismatches 0; Indels 0;

QY 812 CTGCCCCCTTCTCAGCTACTTCG 835
 Db 24 CTGCCCCCTTCTCAGCTACTTCG 1

RESULT 61

AA56003
 ID AAX56003 standard; DNA; 40 BP.

XX

AC AAX56003;

XX

DT 15-JUL-1999 (first entry)

XX

DE Human tumour necrosis factor Apo-3 ligand PCR primer SEQ ID NO:5

XX

KW Human; tumour necrosis factor; Apo-3 ligand; lymphotoxin; apopto

KW NF-kappaB-dependent transcription; JNK/SAPK-dependent response;

KW PCR primer; ss.

XX

OS Synthetic.

OS Homo sapiens.

XX

PN WO9919490-A1.

XX

PD 22-APR-1999.

XX

FF 09-OCT-1998; 98WO-US021407.

XX

PR 10-OCT-1997; 97US-0062037P.

PR 17-DEC-1997; 97US-0069862P.

XX

PA (GETH) GENENTECH INC.

XX

PI Ashkenazi AJ, Marsters SA, Pitti R;

XX

DR WPI; 1999-287982/24.

XX

PT New human Apo3- ligand (a tumor necrosis factor) homologue.

XX

PS Example 2; Page 37; 74pp; English.

XX

CC The present invention describes a human tumour necrosis factor (

CC lymphotoxin homologue designated Apo-3 ligand. Apo-3 ligand has

CC cytostatic activity. Apo-3 ligand can be used to induce apoptosi

CC mammalian cancer cells, to induce NF-kappaB-dependent transcript

CC to induce JNK/SAPK-dependent responses in mammalian cells. The p

CC sequence represents an Apo-3 ligand PCR primer, which is used in

CC example from the present invention

XX

SQ Sequence 40 BP; 9 A; 13 C; 13 G; 5 T; 0 U; 0 Other;

Query Match 1.7%; Score 24; DB 2; Length 40;
 Best Local Similarity 100.0%; Pred.No.1;
 Matches 24; Conservative 0; Mismatches 0; Indels 0;

QY. 244 CGGGCATCGCTGTCGCCGAGGAG 267
 Db 17 CGGGCATCGCTGTCGCCGAGGAG 40

RESULT 62

ABK40355

ID ABK40355 standard; DNA; 23 BP.

XX

AC ABK40355;

XX

DT 15-JUL-2002 (first entry)

XX

DE Probe for gene amplification analysis of human PRO207.

enign tumour; malignant tumour; lymphoid malignancy;
 urological disorder; stromal disorder; blastocoeleic disorder;
 disorder; immune disorder; angiogenic disorder; cytostatic;
 ve; probe; ss.

1.

2000WO-US003565.

99WO-US005028.

99US-0123972P.

99US-0133459P.

99WO-US012252.

99US-0140650P.

99US-0140653P.

99US-0144758P.

99US-0145698P.

99US-0146222P.

99US-0149395P.

99US-0151689P.

99WO-US020111.

99WO-US021090.

99WO-US028313.

99WO-US028301.

99WO-US028634.

2000WO-US000219.

TECH INC.

Goddard A, Godowski PJ, Gurney AL, Hillan KJ;

Pan J, Pitti RM, Roy MA, Smith V, Stone DM;

Wood WI;

567/26.

icletic acids encoding PRO polypeptides, useful for treating
 gnant tumors, leukemias and lymphoid malignancies,
 angiogenic and immunologic disorders.

ige 140; 302pp; English.

vention relates to the isolation of novel human PRO
 AAU86128-AAU86162) and the polynucleotide sequences
 The PRO polypeptides, agonists, antagonists or anti-PRO
 ; useful for treating benign or malignant tumours (e.g.
 bladder, breast, etc), leukaemias and lymphoid
 other disorders such as neuronal, glial, astrocytal,
 glandular, macrophagal, stromal and blastocoeleic disorders,
 immune and angiogenic disorders. The polynucleotide
 also useful in gene therapy. The present sequence
 robe used in the methods of the present invention

; 1 A; 7 C; 7 G; 8 T; 0 U; 0 Other;

1.7%; Score 23; DB 6; Length 23;

arity 100.0%; Pred. No. 3.1;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

TGGGCGCTGTCACGTGTT 1003

TGGGCGCTGTCACGTGTT 23

lard; DNA; 38 BP.

XX 15-JUL-1999 (first entry)
 DT Human tumour necrosis factor Apo-3 ligand PCR primer SEQ ID NO:6.
 DE Human; tumour necrosis factor; Apo-3 ligand; lymphotoxin; apopto;
 KW NF-kappaB-dependent transcription; JNK/SAPK-dependent response; c
 KW PCR primer; ss.
 XX Synthetic.
 OS Homo sapiens.
 XX WO9919490-A1.
 PN 22-APR-1999.
 XX 09-OCT-1998; 98WO-US021407.
 XX 10-OCT-1997; 97US-0062037P.
 PR 17-DEC-1997; 97US-0069862P.
 XX (GETH) GENENTECH INC.
 XX Ashkenazi AJ, Marsters SA, Pitti R;
 XX WPI; 1999-287982/24.
 XX New human Apo3- ligand (a tumor necrosis factor) homologue.
 XX Example 2; Page 37; 74pp; English.

XX The present invention describes a human tumour necrosis factor (T
 CC lymphotoxin homologue designated Apo-3 ligand. Apo-3 ligand has
 CC cytosolic activity. Apo-3 ligand can be used to induce apoptosis
 CC mammalian cancer cells, to induce NF-kappaB-dependent transcripti
 CC to induce JNK/SAPK-dependent responses in mammalian cells. The pr
 CC sequence represents an Apo-3 ligand PCR primer, which is used in
 CC example from the present invention

XX Sequence 38 BP; 9 A; 11 C; 12 G; 6 T; 0 U; 0 Other;

Query Match 1.6%; Score 22; DB 2; Length 38;

Best Local Similarity 100.0%; Pred. No. 8.9;

Matches 22; Conservative 0; Mismatches 0; Indels 0; G

QY 834 CGGACTCTTCCAGGTTCACTGA 855

Db 38 CGGACTCTTCCAGGTTCACTGA 17

RESULT 64

AAU18265

ID AAL18265 standard; cDNA; 223 BP.

XX AAL18265;

XX 07-DEC-2001 (first entry)

DE Human breast cancer expressed polynucleotide 10722.

XX Human; breast cancer; cell marker; cytostatic; ss.

OS Homo sapiens.

XX WO200151628-A2.

PN 19-JUL-2001.

XX 10-JAN-2001; 2001WO-US000798.

XX 14-JAN-2000; 2000US-0176077P.

PR 14-MAR-2000; 2000US-0189167P.

PR 24-MAR-2000; 2000US-0192099P.

2000US-0193480P.
2000US-0205230P.
2000US-0211315P.
2000US-0220534P.

ENNUIUM PREDICTIVE MEDICINE INC.

u Y, Wang Y, Steinmann K;
1856/48.

useful as a marker for the diagnosis of breast cancer.

1912-1913; 3695pp; English.

relates to human breast cancer expressed polynucleotides (ADA07544-ADA26789) and methods of assessing whether a patient is afflicted with breast cancer by examining the correlation between expression of certain markers and the cancerous state of breast cells. The polynucleotides and encoded polypeptides are potential markers for detecting, diagnosing, monitoring, characterising treating and preventing breast cancer. The polynucleotides and encoded polypeptides are also useful for isolating compounds with cytostatic activity.

BP; 46 A; 51 C; 48 G; 78 T; 0 U; 0 Other;

Query Match 1.6%; Score 22; DB 4; Length 223;
Best Local Similarity 100.0%; Pred. No. 8.1;
Matches 0; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

TATTATTTTATTATTATT 1351
|||||
TATTATTTTATTATTATT 119

standard; CDNA; 263 BP.

(first entry)

cancer expressed polynucleotide 1278.

cancer; cell marker; cytostatic; ss.

A2.

2001WO-US000798.

2000US-0176077P.
2000US-0189167P.
2000US-0192099P.
2000US-0193480P.
2000US-0205230P.
2000US-0211315P.
2000US-0220534P.

ENNUIUM PREDICTIVE MEDICINE INC.

u Y, Wang Y, Steinmann K;
1856/48.

useful as a marker for the diagnosis of breast cancer.

278; 3695pp; English.

XX
CC
CC
CC
CC
CC
CC
CC
CC
CC
CC
XX

The invention relates to human breast cancer expressed polynucleotides (ADA07544-ADA26789) and methods of assessing whether a patient is afflicted with breast cancer by examining the correlation between expression of certain markers and the cancerous state of breast cells. The polynucleotides and encoded polypeptides are potential markers for detecting, diagnosing, monitoring, characterising treating and preventing breast cancer. The polynucleotides and encoded polypeptides are also useful for isolating compounds with cytostatic activity.

Sequence 263 BP; 52 A; 63 C; 63 G; 85 T; 0 U; 0 Other;

Query Match 1.6%; Score 22; DB 4; Length 263;
Best Local Similarity 100.0%; Pred. No. 8.1;
Matches 22; Conservative 0; Mismatches 0; Indels 0;

QY 1330 AGATATTTTATTATTATT 1351
|||||

Db 137 AGATATTTTATTATTATT 158
|||||

RESULT 66

ADA02995

ID ADA02995 standard; CDNA; 1005 BP.

AC ADA02995;

XX 06-NOV-2003 (first entry)

Mouse Sept9 carcinoma associated coding sequence, SEQ ID NO:1513
Mouse; murine; carcinoma associated; oncogene; carcinoma; cancer
prostate; lymphoma; leukaemia; cytostatic; gene therapy; drug sc
gene; ss.

Mus sp.

WO2003057146-A2.

17-JUL-2003.

26-DEC-2002; 2002WO-US041414.

26-DEC-2001; 2001US-00035832.

(SAGR-) SAGRES DISCOVERY.

Morris DW;

WPI; 2003-587068/55.

New recombinant nucleic acid encoding carcinoma associated prote
useful for preparing compositions for treating carcinomas.

Claim 1; SEQ ID NO 1513; 245pp; English.

The invention relates to recombinant carcinoma associated (CA) n
acid sequences from mouse and human (ADA01482-ADA03094), and to
recombinant carcinoma associated proteins (CAP) encoded by them.
invention also encompasses expression vectors and host cells com
CA nucleic acid, a polypeptide (especially an antibody) that spe
binds to the protein, and a biochip comprising CA nucleic acid o
fragments thereof. The sequences of the invention were identifie
on oncogenic retroviruses, which insert into the genome of the host
at random. Many of these do not carry transduced host oncogenes
pathogenic trans-acting viral genes, meaning that cancer inciden
direct consequence of the effects of proviral integration into h
protooncogenes. The CA nucleic acid sequences can be used to dia
carcinoma (especially breast cancer, prostate cancer, lymphoma o
leukaemia) or a propensity to carcinoma by determination of the
of a CA gene, or by determination of CA gene expression in part
tissues. CA nucleic acids, proteins and antibodies are also usefi

agents and in screening and evaluating drug candidates. The
 invention represents a specifically claimed murine CA nucleic acid
 sequence. Note: The complete sequence data for this
 invention is available in the printed specification, but was obtained
 directly from WIPO at
 pub/published_pct_sequences.

BP; 285 A; 268 C; 273 G; 179 T; 0 U; 0 Other;

arity 1.6%; Score 22; DB 8; Length 1005;

onservative 0; Mismatches 0; Indels 0; Gaps 0;

CCATTATGAAGTTCATC 448

CCATTATGAAGTTCATC 396

dard; cDNA; 1005 BP.

(first entry)

DNA.

ostatic; gene therapy; vaccine; carcinoma; lymphomas;
 asm; adenocarcinoma; sarcoma; gene.

A2.

2001WO-US051291.

2001US-00798586.

2001US-00004113.

2001US-00052482.

2001US-00997722.

2001US-00034650.

S DISCOVERY.

ngelhard EK;

337/23.

it nucleic acid, useful for treating carcinomas, lymphomas,
 lasm, adenocarcinoma, or sarcomas.

ID NO 561; 2304pp; English.

relates to a novel recombinant nucleic acid comprising a
 sequence selected from any of the 660 sequences fully defined
 in the invention. A polynucleotide of the invention has cytostatic
 activity and may have a use in gene therapy, or in a vaccine. The
 nucleic acids and polypeptides are useful for treating
 g. lymphomas, cancers, neoplasm, adenocarcinoma, and
 present sequence represents a mouse cDNA of the invention.

BP; 285 A; 268 C; 273 G; 179 T; 0 U; 0 Other;

arity 1.6%; Score 22; DB 9; Length 1005;

onservative 0; Mismatches 0; Indels 0; Gaps 0;

CCATTATGAAGTTCATC 448

CCATTATGAAGTTCATC 396

RESULT 68

ADC85475

ID ADC85475 standard; DNA; 1005 BP.

XX

AC ADC85475;

XX

DT 01-JAN-2004 (first entry)

XX

DE Mouse Sept19 coding sequence.

XX

KW Cytostatic; gene therapy; vaccine; cancer; carcinoma-associated;
 KW secreted; transmembrane; intracellular; ds.

XX

OS Mus sp.

XX

PN WO2003045230-A2.

XX

PD 05-JUN-2003.

XX

PF 02-DEC-2002; 2002WO-US038582.

XX

PR 30-NOV-2001; 2001US-00997722.

XX

PA (SAGR-) SAGRES DISCOVERY.

XX

PI Morris DW, Engelhard EK;

XX

DR WPI; 2003-513603/48.

XX

PT New recombinant nucleic acid comprising a nucleotide sequence of
 PT the carcinoma-associated (CA) genes, useful for screening for dru
 PT candidates for diagnosing or treating carcinomas.

XX

PS Claim 1; SEQ ID NO 261; 983pp; English.

XX

CC The invention relates to a recombinant nucleic acid comprising a
 CC nucleotide sequence selected from any of the fully defined carcin
 CC associated (CA) genes from the 50 tables given in the specificati
 CC CA proteins are secreted, transmembrane or intracellular proteins
 CC recombinant nucleic acids are useful for screening for drug candi
 CC for diagnosing or treating carcinomas. Sequences given in ADC8521
 CC ADC85514 represent CA genes of the invention.

XX

SQ Sequence 1005 BP; 285 A; 268 C; 273 G; 179 T; 0 U; 0 Other;

Query Match

Best Local Similarity 1.6%; Score 22; DB 9; Length 1005;

Matches 22; Conservative 0; Mismatches 0; Indels 0; G

QY 427 GCAGCCCATTCAGTTCATC 448

Db 375 GCAGCCCATTCAGTTCATC 396

RESULT 69

AAS84907

ID AAS84907 standard; cDNA; 1778 BP.

XX

AC AAS84907;

XX

DT 13-FEB-2002 (first entry)

XX

DE DNA encoding novel human diagnostic protein #20711.

XX

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; s;

XX

OS Homo sapiens.

XX

PN WO200175067-A2.

XX

2001WO-US008631.
2000US-00540217.
2000US-00649167.
Q INC.
Liu C, Tang YT;
9362/73.
0720.
polynucleotide and encoded polypeptides, useful in forensics, gene mapping, identification of mutations for genetic disorders or other traits and to assess
ID NO 20711; 103pp; English.
n relates to isolated polynucleotide (I) and polypeptide (II) I) is useful as hybridisation probes, polymerase chain R) primers, oligomers, and for chromosome and gene mapping, binant production of (II). The polynucleotides are also used as expressed sequence tags for identifying expressed (II) or to treat disease states involving (II). (II) is generating antibodies against it, detecting or quantitating a in tissue, as molecular weight markers and as a food (II) and its binding partners are useful in medical imaging errant protein expression or biological activity. The and polynucleotide sequences have applications in forensics, gene mapping, identification of mutations for genetic disorders or other traits to assess biodiversity ce other types of data and products dependent on DNA and sequences. AAS64197-AAS94564 represent novel human diagnostic ces of the invention. Note: The sequence data for this at appear in the printed specification, but was obtained in ormat directly from WIPO at /pub/published_pct_sequences
3 BP; 472 A; 408 C; 371 G; 527 T; 0 U; 0 Other;
arity 100.0%; Pred. No. 7.3;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
3GATGGGGGGGGCGGTGAGG 84
3GATGGGGGGGGCGGTGAGG 205
idard; cDNA; 2942 BP.
(first entry)
carcinoma associated cDNA, SEQ ID NO:1512.
; carcinoma associated; oncogene; carcinoma; cancer; breast; mphoma; leukaemia; cytostatic; gene therapy; drug screening;
-A2.
26-DEC-2002; 2002WO-US041414.
26-DEC-2001; 2001US-00035832.
(SAGR-) SAGRES DISCOVERY.
Morris DW;
WPI; 2003-587068/55.
New recombinant nucleic acid encoding carcinoma associated prote useful for preparing compositions for treating carcinomas.
Claim 1; SEQ ID NO 1512; 245pp; English.
The invention relates to recombinant carcinoma associated (CA) n acid sequences from mouse and human (ADA01482-ADA03094), and to recombinant carcinoma associated proteins (CAP) encoded by them. invention also encompasses expression vectors and host cells com CA nucleic acid, a polypeptide (especially an antibody) that spe binds to the protein, and a biochip comprising CA nucleic acid o fragments thereof. The sequences of the invention were identifie oncogenic retroviruses, which insert into the genome of the host at random. Many of these do not carry transduced host oncogenes pathogenic trans-acting viral genes, meaning that cancer inciden direct consequence of the effects of proviral integration into h protooncogenes. The CA nucleic acid sequences can be used to dia carcinoma (especially breast cancer, prostate cancer, lymphoma o leukaemia) or a propensity to carcinoma by determination of the of a CA gene, or by determination of CA gene expression in parti tissues. CA nucleic acids, proteins and antibodies are also usef therapeutic agents and in screening and evaluating drug candidat present sequence represents a specifically claimed murine CA nuc sequence of the invention. Note: The complete sequence data for patent did not form part of the printed specification, but was o in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.
SQ Sequence 2942 BP; 702 A; 837 C; 780 G; 623 T; 0 U; 0 Other;
Query Match 1.6%; Score 22; DB 8; Length 2942;
Best Local Similarity 100.0%; Pred. No. 7.2;
Matches 22; Conservative 0; Mismatches 0; Indels 0;
QY 427 GCAGCCCATTTAGAGTTCATC 448
Db 552 GCAGCCCATTTAGAGTTCATC 573
RESULT 71
ADB72732
ID ADB72732 standard; mRNA; 2942 BP.
XX
AC ADB72732;
XX
DT 04-DEC-2003 (first entry)
XX
DE Mouse Sept9 mRNA.
XX
KW mouse; ss; cytostatic; gene therapy; vaccine; carcinoma; lymphoma cancer; neoplasm; adenocarcinoma; sarcoma.
XX
OS Mus sp.
XX
PN WO2003008583-A2.
XX
PD 30-JAN-2003.
XX
PF 26-DEC-2001; 2001WO-US051291.
XX
PR 02-MAR-2001; 2001US-00798586.
PR 23-OCT-2001; 2001US-00004113.

2001US-00052482.
2001US-00997722.
2001US-00034650.

S DISCOVERY.

ngelhard EK;

337/23.

nt nucleic acid, useful for treating carcinomas, lymphomas,
lasm, adenocarcinoma, or sarcomas.

ID NO 560; 2304pp; English.

relates to a novel recombinant nucleic acid comprising a
quence selected from any of the 660 sequences fully defined
ication. A polynucleotide of the invention has cytostatic
may have a use in gene therapy, or in a vaccine. The
nucleic acids and polypeptides are useful for treating
.g. lymphomas, cancers, neoplasm, adenocarcinoma, and
present sequence represents a mouse mRNA of the invention.

BP; 702 A; 837 C; 780 G; 623 T; 0 U; 0 Other;

arity 1.6%; Score 22; DB 9; Length 2942;

onservative 0; Mismatches 0; Indels 0; Gaps 0;

CCCATTTATGAAGTTCATC 448

CCCATTTATGAAGTTCATC 573

iard; DNA; 2942 BP.

(first entry)

RNA sequence.

ne therapy; vaccine; cancer; carcinoma-associated gene; CA;
membrane; intracellular; ds.

12.

2002WO-US038582.

2001US-00997722.

S DISCOVERY.

ngelhard EK;

103/48.

it nucleic acid comprising a nucleotide sequence of any of
associated (CA) genes, useful for screening for drug
diagnosing or treating carcinomas.

ID NO 260; 983pp; English.

relates to a recombinant nucleic acid comprising a
quence selected from any of the fully defined carcinoma-
genes from the 50 tables given in the specification. The
e secreted, transmembrane or intracellular proteins. The

CC recombinant nucleic acids are useful for screening for drug candi
for diagnosing or treating carcinomas. Sequences given in ADC8521
CC ADC85514 represent CA genes of the invention.

XX Sequence 2942 BP; 702 A; 837 C; 780 G; 623 T; 0 U; 0 Other;

Query Match

Best Local Similarity 1.6%; Score 22; DB 9; Length 2942;

Matches 22; Conservative 0; Mismatches 0; Indels 0; C

Oy 427 GCAGCCCATTTATGAAGTTCATC 448

Db 552 GCAGCCCATTTATGAAGTTCATC 573

RESULT 73

ADA02993

ID ADA02993 standard; DNA; 50295 BP.

XX ADA02993;

XX ADA02993;

DT 06-NOV-2003 (first entry)

XX Mouse Sept9 carcinoma associated gene, SEQ ID NO:1511.

XX Mouse; murine; carcinoma associated; oncogene; carcinoma; cancer;
prostate; lymphoma; leukaemia; cytostatic; gene therapy; drug scr
gene; ds.

XX Mus sp.

XX WO2003057146-A2.

XX 17-JUL-2003.

XX 26-DEC-2002; 2002WO-US041414.

XX 26-DEC-2001; 2001US-00035832.

XX (SAGR-) SAGRES DISCOVERY.

XX Morris DW;

XX WPI; 2003-587068/55.

XX New recombinant nucleic acid encoding carcinoma associated protei
useful for preparing compositions for treating carcinomas.
Claim 1; SEQ ID NO 1511; 245pp; English.

The invention relates to recombinant carcinoma associated (CA) nu
acid sequences from mouse and human (ADA01482-ADA03094), and to
recombinant carcinoma associated proteins (CAP) encoded by them.
invention also encompasses expression vectors and host cells comp.
CA nucleic acid, a polypeptide (especially an antibody) that spec
binds to the protein, and a biochip comprising CA nucleic acid or
fragments thereof. The sequences of the invention were identified
oncogenic retroviruses, which insert into the genome of the host
at random. Many of these do not carry transduced host oncogenes o
pathogenic trans-acting viral genes, meaning that cancer incidence
direct consequence of the effects of proviral integration into ho
protooncogenes. The CA nucleic acid sequences can be used to diagn
carcinoma (especially breast cancer, prostate cancer, lymphoma or
leukaemia) or a propensity to carcinoma by determination of the se
of a CA gene, or by determination of CA gene expression in partic
tissues. CA nucleic acids, proteins and antibodies are also useful
therapeutic agents and in screening and evaluating drug candidate
present sequence represents a specifically claimed murine CA nucle
sequence of the invention. Note: The complete sequence data for th
patent did not form part of the printed specification, but was obt
in electronic format directly from WIPO at
ftp.wipo.int/pub/published_pct_sequences.

XX

06:25:14 2004

us-09-245-198a-3.oligo.rng

95 BP; 10891 A; 12938 C; 13611 G; 12237 T; 0 U; 618 Other;
1.6%; Score 22; DB 8; Length 50295;
arity 100.0%; Pred. No. 6.2;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
GCCATTATGAAGTTTCATC 448
|||||
GCCATTATGAAGTTTCATC 31093

ndard; DNA; 50295 BP.

(first entry)
gene.
ytostatic; gene therapy; vaccine; carcinoma; lymphomas;
laem; adenocarcinoma; sarcoma; gene.
-A2.
2001WO-US051291.
2001US-00798586.
2001US-00004113.
2001US-00052482.
2001US-00997722.
2001US-00034650.
ES DISCOVERY.
Engelhard EK;
3337/23.
ant nucleic acid, useful for treating carcinomas, lymphomas,
plasm, adenocarcinoma, or sarcomas.
ID NO 559; 2304pp; English.
relates to a novel recombinant nucleic acid comprising a
sequence selected from any of the 660 sequences fully defined
ication. A polynucleotide of the invention has cytostatic
i may have a use in gene therapy, or in a vaccine. The
nucleic acids and polypeptides are useful for treating
g. lymphomas, cancers, neoplasm, adenocarcinoma, and
; present sequence represents a mouse gene of the invention.
95 BP; 10891 A; 12938 C; 13611 G; 12237 T; 0 U; 618 Other;
1.6%; Score 22; DB 9; Length 50295;
arity 100.0%; Pred. No. 6.2;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
GCCATTATGAAGTTTCATC 448
|||||
GCCATTATGAAGTTTCATC 31093

ndard; DNA; 50295 BP.

XX 01-JAN-2004 (first entry)
DT Mouse Sept19 genomic sequence.
DE
XX Cytostatic; gene therapy; vaccine; cancer; carcinoma-associated
KW secreted; transmembrane; intracellular; ds.
XX
OS Mus sp.
XX WO2003045230-A2.
PN
XX 05-JUN-2003.
PD
XX 02-DEC-2002; 2002WO-US038582.
PF
XX 30-NOV-2001; 2001US-00997722.
PR
XX (SAGR-) SAGRES DISCOVERY.
PA
XX Morris DW, Engelhard EK;
PI WPI; 2003-513603/48.
DR
XX New recombinant nucleic acid comprising a nucleotide sequence of
PT the carcinoma-associated (CA) genes, useful for screening for dr
PT candidates for diagnosing or treating carcinomas.
XX
PS Claim 1; SEQ ID NO 259; 983pp; English.
XX
CC The invention relates to a recombinant nucleic acid comprising a
CC nucleotide sequence selected from any of the fully defined carci
CC associated (CA) genes from the 50 tables given in the specificat
CC CA proteins are secreted, transmembrane or intracellular protein
CC recombinant nucleic acids are useful for screening for drug cand
CC for diagnosing or treating carcinomas. Sequences given in ADC852
CC ADC8514 represent CA genes of the invention.
XX
SQ Sequence 50295 BP; 10891 A; 12938 C; 13611 G; 12237 T; 0 U; 618
Query Match 1.6%; Score 22; DB 9; Length 50295;
Best Local Similarity 100.0%; Pred. No. 6.2;
Matches 22; Conservative 0; Mismatches 0; Indels 0;
QY 427 GCAGCCCATTTATGAAGTTTCATC 448
Db 31072 GCAGCCCATTTATGAAGTTTCATC 31093

Search completed: April 8, 2004, 21:02:41
Job time : 662 secs

GenCore version 5.1.6

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in search, using sw model

ril 7, 2004, 17:54:48 ; Search time 20 Seconds
(without alignments)
1365.920 Million cell updates/sec

-09-245-198a-4

MSLLDFEISARRLPLPRSLG.....PWAHLKAAPFLTYRGLFQVH 284

pop 60.0 , Gapext 60.0

3366 seqs, 96191526 residues

ts satisfying chosen parameters: 283366

3th: 0

3th: 20000000000

isting first 100 summaries

IR 78:*

Pir1:*

Pir2:*

Pir3:*

Pir4:*

the number of results predicted by chance to have a
score than or equal to the score of the result being printed,
and by analysis of the total score distribution.

SUMMARIES

seq	len	db	id	Description
1.5	111	2	A85866	hypothetical prote
1.2	733	2	S78376	photosystem I p700
1.8	58	2	A58208	protamine I, 1 - pa
1.8	143	2	G84168	hypothetical prote
1.8	197	2	E72374	hypothetical prote
1.8	220	2	AG3547	bicyclomycin resis
1.8	278	2	D83080	hypothetical prote
1.8	339	2	C71132	hypothetical prote
1.8	372	2	H70813	hypothetical prote
1.8	379	2	E64300	probable cysteine
1.8	381	2	AH3041	formate dehydrogen
1.8	387	2	D84885	conserved hypothet
1.8	397	2	D98244	hypothetical prote
1.8	422	1	A60503	hypothetical prote
1.8	443	2	T17220	sperm-binding glyc
1.8	455	2	AC0347	hypothetical prote
1.8	471	2	A75267	probable membrane
1.8	576	2	E64186	probable transport
1.8	586	2	A41125	probable ATP-bind
1.5	45	2	D58208	gamma-glutamyltran
1.5	50	2	S22582	protamine II-3 - p
1.5	58	2	S34045	protamine 1 - Sagu
1.5	86	2	F87604	protamine - North
1.5	102	2	F87993	hypothetical prote
1.5	115	2	PH1560	protein ZC334.3 [i
1.5	115	2	H83201	Ig heavy chain V r
1.5	118	1	IEEC5B	conserved hypothet
1.5	118	2	AE1753	hypothetical prote
1.5	123	2	AH2707	Orf51 bacteriophag
1.5	123	2	AH2707	conserved hypothet

30	7	2.5	125	2	T27519	hypothetical
31	7	2.5	131	2	I52290	interleukin
32	7	2.5	131	2	E30552	T-cell act;
33	7	2.5	146	2	T37116	probable t;
34	7	2.5	147	2	A71217	hypothetical
35	7	2.5	150	2	T08734	hypothetical
36	7	2.5	157	2	S31078	seed aller
37	7	2.5	157	2	T02664	allergen -
38	7	2.5	157	2	A75567	conserved l
39	7	2.5	157	2	E75530	hypothetical
40	7	2.5	160	2	S59925	allergen R
41	7	2.5	161	1	DNEC17	outer membr
42	7	2.5	161	2	D90651	histone-li
43	7	2.5	161	2	D85502	hypothetical
44	7	2.5	162	2	T24937	hypothetical
45	7	2.5	162	2	T31173	hypothetical
46	7	2.5	164	2	S76920	hypothetical
47	7	2.5	170	2	S44789	D2007.4 prc
48	7	2.5	174	2	D87638	transcripti
49	7	2.5	180	1	LGST	beta-lactog
50	7	2.5	180	1	LGSH	beta-lactog
51	7	2.5	181	2	B60738	insulin-li
52	7	2.5	187	2	G85343	phospholipa
53	7	2.5	206	2	S72567	hypothetical
54	7	2.5	230	2	AH0692	conserved h
55	7	2.5	231	2	B64920	probable me
56	7	2.5	231	2	E90921	hypothetical
57	7	2.5	231	2	A85770	hypothetical
58	7	2.5	231	2	C86665	amino acid
59	7	2.5	233	2	S60767	ribonucleas
60	7	2.5	233	2	AB0273	probable me
61	7	2.5	234	2	G85098	H+-transport
62	7	2.5	235	2	I64174	probable so
63	7	2.5	236	2	A86387	probable cy
64	7	2.5	238	2	H70866	hypothetical
65	7	2.5	240	2	G83208	conserved h
66	7	2.5	243	2	C64124	molybdopter
67	7	2.5	243	2	T25942	hypothetical
68	7	2.5	244	2	A46066	lymphotoxin
69	7	2.5	249	2	B32352	molybdopter
70	7	2.5	249	2	C85592	molybdopter
71	7	2.5	249	2	H90741	molybdopter
72	7	2.5	249	2	AI0602	molybdopter
73	7	2.5	252	2	AI3154	IS21 family
74	7	2.5	252	2	H98132	1stB protei
75	7	2.5	254	2	F82439	molybdopter
76	7	2.5	255	2	AB0182	molybdopter
77	7	2.5	258	2	AI0566	hydroxypyru
78	7	2.5	258	2	G83101	probable ac
79	7	2.5	260	2	AG0007	lipopolysac
80	7	2.5	262	2	F71801	flagellar b
81	7	2.5	262	2	A64718	flagellar b
82	7	2.5	277	2	G75518	probable bel
83	7	2.5	293	2	A97396	hypothetical
84	7	2.5	300	2	AB2614	hypothetical
85	7	2.5	303	2	E87280	hard protei
86	7	2.5	307	2	S23780	nucleic aci
87	7	2.5	307	2	A83466	maltose tra
88	7	2.5	308	2	D95932	probable su
89	7	2.5	310	2	D70328	histidine k
90	7	2.5	312	2	A31846	130K paracr
91	7	2.5	313	2	C75208	sugar trans
92	7	2.5	314	1	A46489	pan-epitheli
93	7	2.5	316	2	C82561	drug tolera
94	7	2.5	318	2	AD3295	sodium/bile
95	7	2.5	319	2	C95927	probable su
96	7	2.5	321	2	B72604	hypothetical
97	7	2.5	325	2	C36718	pyruvate def
98	7	2.5	329	2	T18619	hypothetical
99	7	2.5	333	2	B90172	conserved h
100	7	2.5	343	2	A43577	regulatory f

ALIGNMENTS

ein 23516 [imported] - Escherichia coli (strain O157:H7, substrain EDL933)
 ichia coli
 101 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001
 66
 unkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew
 ck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
 33, 2001
 equence of enterohemorrhagic Escherichia coli O157:H7.
 r: A85480; MUID:21074935; PMID:11206551
 66
 nary
 DNA

<STO>
 s: GB:AE005174; NID:g12516604; PIDN:AAG57389.1; GSPDB:GN00145; UWGP:Z35
 urce: strain O157:H7, substrain EDL933

3.5%; Score 10; DB 2; Length 111;
 larity 100.0%; Pred. No. 0.074;
 Conservative 0; Mismatches 0; Indels 0; Gaps 0;

ALACGL 72
 |||||
 ALACGL 49

0 apoprotein A2 - Odontella sinensis chloroplast
 plast Odontella sinensis
 98 #sequence_revision 26-Feb-1998 #text_change 20-Jun-2000
 76
 Stoebe, B.; Schaffran, I.; Kroth-Pancic, P.; Freier, U.
 Rep. 13, 336-342, 1995
 oplast Genome of a chlorophyll a+c- containing Alga, Odontella sinensis
 c: S78238
 76

nary; nucleic acid sequence not shown; translation not shown
 DNA

<KOW>
 i: EMBL:267753; NID:g1185127; PIDN:CAA91749.1; PID:g1185266
 stide sequence was submitted to the EMBL Data Library, November 1995

ast
 rosystem I p700 apoprotein
 plast; electron transfer; membrane protein; membrane-associated complex

3.2%; Score 9; DB 2; Length 733;
 arity 100.0%; Pred. No. 3.3;
 onservative 0; Mismatches 0; Indels 0; Gaps 0;

LAACLG 71
 |||||
 LAACLG 341

inted turtle
 ys picta (painted turtle)
 6 #sequence_revision 08-Nov-1996 #text_change 07-May-1999
 8
 nsky, H.E.; Elsev, R.M.; Wright, C.L.; Rice, P.; Bell, J.E.; Sharp, D.
 , 23547-23557, 1996
 s of reptiles.
 : A58208; MUID:96394458; PMID:8798564

A;Accession: A58208
 A;Status: preliminary
 A;Molecule type: protein
 A;Residues: 1-58 <HUN>
 C;Superfamily: sperm histone

Query Match 2.8%; Score 8; DB 2; Length 58;
 Best Local Similarity 100.0%; Pred. No. 3.5;
 Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 42 ORRRGRGG 49
 |||||
 Db 35 ORRRGRGG 42

RESULT 4

G84168
 hypothetical protein Vng0080h [imported] - Halobacterium sp. NRC-1
 C;Species: Halobacterium sp. NRC-1
 C;Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
 C;Accession: G84168

R;Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shu
 ; Leithauser, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Ma
 Jung, K.H.; Alam, M.; Freitas, T.
 Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000
 A;Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt,
 A;Title: Genome sequence of Halobacterium species NRC-1.
 A;Reference number: A84160; MUID:20504483; PMID:11018950

A;Accession: G84168

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-143 <STO>

A;Cross-references: GB:AE004437; NID:g10579733; PIDN:AAG18715.1; GSPD
 C;Genetics:

A;Gene: VNG0080H

Query Match 2.8%; Score 8; DB 2; Length 143;
 Best Local Similarity 100.0%; Pred. No. 7.6;
 Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 61 LGLGLALA 68
 |||||
 Db 55 LGLGLALA 62

RESULT 5

E72374

hypothetical protein - Thermotoga maritima (strain MSB8)

C;Species: Thermotoga maritima

C;Date: 11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 21-Jun-1999
 C;Accession: E72374

R;Nelson, K.E.; Clayton, R.A.; Gill, S.R.; Gwinn, M.L.; Dodson, R.J.;
 Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.;
 C.M.

Nature 399, 323-329, 1999

A;Title: Evidence for lateral gene transfer between Archaea and Bacter

A;Reference number: A72200; MUID:99287316; PMID:10360571

A;Accession: E72374

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-197 <ARN>

A;Cross-references: GB:AE001724; GB:AE000512; NID:g4980966; PIDN:AAD35
 A;Experimental source: strain MSB8

C;Genetics:

A;Gene: TM0469

Query Match 2.8%; Score 8; DB 2; Length 197;
 Best Local Similarity 100.0%; Pred. No. 10;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 2 SLDDFEIS 9
 |||||
 Db 135 SLDDFEIS 142

A;Accession: C71132
 A;Status: preliminary; nucleic acid sequence not shown; translation nc
 A;Molecule type: DNA
 A;Residues: 1-339 <RAW>
 A;Cross-references: GB:AP000003; NID:g3236130; PIDN:BAA29917.1; PID:g3
 A;Experimental source: strain OT3
 A;Note: this accession replaces an interim accession for a sequence re
 C;Genetics:
 A;Gene: PH0824
 C;Superfamily: conserved hypothetical protein MTH900

Query Match 2.8%; Score 8; DB 2; Length 339;
 Best Local Similarity 100.0%; Pred. No. 16;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 70 LGLLLAVV 77
 Db 111 LGLLLAVV 118

RESULT 9
 H70813
 probable cysteine synthase - Mycobacterium tuberculosis (strain H37RV)
 C;Species: Mycobacterium tuberculosis
 C;Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 20-Jun
 C;Accession: H70813
 R;Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Har
 ; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamli
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squ
 Nature 393, 537-544, 1998
 A;Authors: Sgares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barre
 A;Title: Deciphering the biology of Mycobacterium tuberculosis from th
 A;Reference number: A70500; MUID:98295987; PMID:9634230
 A;Accession: H70813
 A;Status: preliminary; nucleic acid sequence not shown; translation no
 A;Molecule type: DNA
 A;Residues: 1-372 <COL>
 A;Cross-references: GB:AL020004; GB:AL123456; NID:g3261550; PIDN:CAA17
 A;Experimental source: strain H37RV
 C;Genetics:
 A;Gene: cysM3
 C;Superfamily: threonine dehydratase

Query Match 2.8%; Score 8; DB 2; Length 372;
 Best Local Similarity 100.0%; Pred. No. 17;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 61 LGLGLALA 68
 Db 96 LGLGLALA 103

RESULT 10
 E64300
 formate dehydrogenase (EC 1.2.1.2) beta chain - Methanococcus jannaschi
 C;Species: Methanococcus jannaschi
 C;Date: 13-Sep-1996 #sequence_revision 13-Sep-1996 #text_change 21-Jul-
 C;Accession: E64300
 R;Bult, C.J.; White, O.; Olsen, G.J.; Zhou, L.; Fleischmann, R.D.; Sutt
 ; Reich, C.I.; Overbeek, R.; Kirkness, E.F.; Weinstock, K.G.; Merrick,
 rson, J.D.; Sadow, P.W.; Hanna, M.C.; Cotton, M.D.; Roberts, K.M.; Hurs
 Science 273, 1058-1073, 1996
 A;Authors: Kaine, B.P.; Borodovsky, M.; Klenk, H.P.; Fraser, C.M.; Smit
 A;Title: Complete genome sequence of the methanogenic archaeon, Methano
 A;Reference number: A64300; MUID:96337999; PMID:8688087
 A;Accession: E64300
 A;Status: preliminary; nucleic acid sequence not shown; translation not
 A;Molecule type: DNA
 A;Residues: 1-379 <BU>
 A;Cross-references: GB:U67459; GB:L77117; NID:g2826236; PIDN:AAB97986.1
 C;Genetics:
 A;Map position: REV7250-6111
 C;Superfamily: formate dehydrogenase chain B; ferredoxin 2 [4Fe-4S] homo

tance protein [imported] - Brucella melitensis (strain 16M)
 a melitensis
 2 #sequence_revision 01-Feb-2002 #text_change 01-Feb-2002
 7
 ; Kapatal, V.; Redkar, R.J.; Patra, G.; Mujer, C.; Los, T.; Ivanova,
 sman, E.; Selkov, E.; Elzer, P.H.; Hagius, S.; O'Callaghan, D.; Letess
 Sci. U.S.A. 99, 443-448, 2002
 a sequence of the facultative intracellular pathogen Brucella melitens
 ; AD3252; PMID:11756688
 ;
 ;ry
 ;NA
 <KUR>
 ; GB:AE008918; PIDN:AAL53546.1; PID:g17984455; GSPDB:GN00191
 ce: strain 16M

2.8%; Score 8; DB 2; Length 220;
 rity 100.0%; Pred. No. 11;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 LAL 61
 ||||
 LAL 139

n PA4521 [imported] - Pseudomonas aeruginosa (strain PA01)
 nas aeruginosa
 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
 m, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.; Br
 ; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lim,
 M.V.
 , 2000
 enome sequence of Pseudomonas aeruginosa PA01, an opportunistic patho
 A82950; MUID:20437337; PMID:10984043
 ry
 A
 STO>
 GB:AE004866; GB:AE004091; NID:g9950760; PIDN:AAG07909.1; GSPDB:GN001
 ce: strain PA01

2.8%; Score 8; DB 2; Length 278;
 rity 100.0%; Pred. No. 13;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 AVW 77
 ||||
 AVW 54

n PH0824 - Pyrococcus horikoshii
 us horikoshii
 #sequence_revision 14-Aug-1998 #text_change 20-Jun-2000
 Sawada, M.; Horikawa, H.; Haikawa, Y.; Hino, Y.; Yamamoto, S.; Sekin
 nahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; Kushida, N.; Oguchi
 1998
 equence and gene organization of the genome of a hyper-thermophilic a
 A71000; MUID:98344137; PMID:9679194

reductase
ferredoxin 2[4Fe-4S] homology <FER>
2.8%; Score 8; DB 2; Length 379;
arity 100.0%; Pred. No. 17;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
3VAL 224
|||||
3VAL 42

ical protein Atu3948 [imported] - Agrobacterium tumefaciens (strain C58)
2 #sequence_revision 11-Jan-2002 #text_change 18-Nov-2002
11
bal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo, L.
W.; Grant, C.; Guenther, D.; Kutayin, T.; Levy, R.; Li, M.; McClellan,
P.; Zhang, S.
2323, 2001
; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm,
ne of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
C: AB2577; MUID:21608550; PMID:11743193
11
ary
DNA
<KUR>
3: GB:AE008689; PIDN:AAL44750.1; PID:G17742385; GSPDB:GN00187
rce: strain C58 (Dupont)

near chromosome
2.8%; Score 8; DB 2; Length 381;
arity 100.0%; Pred. No. 17;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
PGSS 258
|||||
PGSS 332

in At2g45000 [imported] - Arabidopsis thaliana
psis thaliana (mouse-ear cress)
11 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
15
; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;
t, K.S.; Cronin, L.A.; Shen, M.; VanAken, S.E.; Umayam, L.; Tallon, L.
W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, J.
8, 1999
and analysis of chromosome 2 of the plant Arabidopsis thaliana.
; AB4420; MUID:20083487; PMID:10617197
5
ary
NA
<STO>
1: GB:AE002093; NID:G4895250; PIDN:AAD32835.1; GSPDB:GN00139

2.8%; Score 8; DB 2; Length 387;
arity 100.0%; Pred. No. 18;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
AEED 100
|||||
AEED 377

RESULT 13
D98244
hypothetical protein AGR_L_1808 [imported] - Agrobacterium tumefaciens
C:Species: Agrobacterium tumefaciens
C:Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 18-Nov-2002
C:Accession: D98244
R:Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Qu
A.; Liu, F.; Wollam, C.; Allinger, M.; Dougherty, D.; Scott, C.; Lappa
Science 294, 2323-2328, 2001
A:Title: Genome Sequence of the Plant Pathogen and Biotechnology Agen
A:Reference number: A97359; MUID:21608551; PMID:11743194
A:Accession: D98244
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-397 <KUR>
A:Cross-references: GB:AE007870; PIDN:AAK89478.1; PID:G15159347; GSPD
C:Genetics:
A:Gene: AGR_L_1808
A:Map position: linear chromosome

Query Match 2.8%; Score 8; DB 2; Length 397;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 8; Conservative 0; Mismatches 0; Indels 0;
QY 251 LALRPGSS 258
|||||
Db 341 LALRPGSS 348

RESULT 14
A60503
sperm-binding glycoprotein ZP3 precursor - Golden hamster
N:Alternate names: sperm receptor; zona pellucida glycoprotein ZP3
C:Species: Mesocricetus auratus (golden hamster)
C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999
C:Accession: A60503
R:Kinloch, R.A.; Ruiz-Seiler, B.; Wasserman, P.M.
Dev. Biol. 142, 414-421, 1990
A:Title: Genomic organization and polypeptide primary structure of zo
A:Reference number: A60503; MUID:91078540; PMID:2257975
A:Accession: A60503
A:Molecule type: DNA
A:Residues: 1-422 <KIN>
A:Cross-references: GB:M63629
A:Note: the authors translated the codon CAA for residue 251 as Glu, z
C:Comment: This sulfated glycoprotein in the zona pellucida of the ooc
C:Superfamily: sperm-binding glycoprotein ZP3; ZP domain homology
C:Keywords: glycoprotein; oocyte
F:45-300/Domain: ZP domain homology <ZPH>

Query Match 2.8%; Score 8; DB 1; Length 422;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 0; Indels 0;
QY 59 LALGLGLA 66
|||||
Db 386 LALGLGLA 393

RESULT 15
T17220
hypothetical protein DKFZp5660011.1 - human
C:Species: Homo sapiens (man)
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 02-Jun-2001
C:Accession: T17220
R:Blum, H.; Baerach, S.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
submitted to the Protein Sequence Database, September 1999
A:Reference number: Z18725
A:Accession: T17220
A:Status: preliminary
A:Molecule type: mRNA

```

<BLU>
: EMBL:AL117414
rce: fetal kidney; clone DKFP5660011

11.1
na-glutamyltransferase
      2.8%; Score 8; DB 2; Length 443;
arity 100.0%; Pred. No. 20;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

LALA 68
|||||
LALA 21

protein yegB [imported] - Yersinia pestis (strain CO92)
a pestis
1 #sequence_revision 02-Nov-2001 #text_change 17-May-2002
en, B.W.; Thomson, N.R.; Titball, R.W.; Holden, M.T.G.; Prentice, M.B.;
: Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis, P.; Dougan, G.;
: K.; Simmonds, M.; Skelton, J.; Stevens, R.; Whitehead, S.; Barrell,
7, 2001
ence of Yersinia pestis, the causative agent of plague.
: AB0001; MUID:21470413; PMID:11586360
ry
JA
:KUR>
: GB:AL590842; PIDN:CAC92102.1; PID:g15980820; GSPDB:GN00175

:drug-efflux transporter
      2.8%; Score 8; DB 2; Length 465;
arity 100.0%; Pred. No. 21;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

VSL 79
|||||
VSL 342

protein - Deinococcus radiodurans (strain R1)
#sequence_revision 03-Dec-1999 #text_change 31-Mar-2000
J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;
: thevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; Ma
577, 1999
ence of the radioresistant bacterium Deinococcus radiodurans R1.
A75250; MUID:20036896; PMID:10567266

ry
A
WHI>
GB:AE002079; GB:AE000513; NID:g6460315; PIDN:AAF12043.1; PID:g646032
ce: strain R1

      2.8%; Score 8; DB 2; Length 471;
arity 100.0%; Pred. No. 21;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

LAL 67

```

```

Db      366 ALGLGLAL 373
|||||||

RESULT 18
E64186
probable ATP-binding transport protein H1156 - Haemophilus influenzae
C:Species: Haemophilus influenzae
C:Date: 18-Aug-1995 #sequence_revision 18-Aug-1995 #text_change 02-Feb
C:Accession: E64186
R;Fleischmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness,
: Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley
: D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Ge
Science 269, 496-512, 1995
A:Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smi
A:Title: Whole-genome random sequencing and assembly of Haemophilus in
A:Reference number: A64000; MUID:95350630; PMID:7542800
A:Accession: E64186
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-576 <TIGR>
A:CROSS-references: GB:U32795; GB:I42023; NID:g1574708; PIDN:AA22811.
C:Superfamily: unassigned ATP-binding cassette proteins; ATP-binding c
C:Keywords: ATP; nucleotide binding; P-loop
F:355-550/Domain: ATP-binding cassette homology <ABC>
F:372-379/Region: nucleotide-binding motif A (P-loop)

Query Match      2.8%; Score 8; DB 2; Length 576;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY      58 PLALGLGL 65
|||||||

Db      159 PLALGLGL 166
|||||||

RESULT 19
A41125
gamma-glutamyltransferase (EC 2.3.2.2) related protein - human
N:Alternate names: gamma-glutamyltransferase-like activity 1; GGT-REL
C:Species: Homo sapiens (man)
C:Date: 27-Mar-1992 #sequence_revision 27-Mar-1992 #text_change 18-Jun
C:Accession: A41125
R;Heisterkamp, N.; Rajpert-De Meyts, E.; Uribe, L.; Forman, H.J.; Grof.
proc. Natl. Acad. Sci. U.S.A. 88, 6303-6307, 1991
A:Title: Identification of a human gamma-glutamyl cleaving enzyme rela
A:Reference number: A41125; MUID:91296809; PMID:1676842
A:Accession: A41125
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-586 <HEI>
A:CROSS-references: GB:M64099; NID:g183141; PIDN:AA58503.1; PID:g1831.
C:Genetics:
A:Gene: GDB:GGTAL1; GGT-REL
A:CROSS-references: GDB:134033
C:Superfamily: gamma-glutamyltransferase
C:Keywords: aminocyltransferase; glycoprotein; transmembrane protein

Query Match      2.8%; Score 8; DB 2; Length 586;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY      61 LGLGLALA 68
|||||||

Db      14 LGLGLALA 21
|||||||

RESULT 20
D58208
protamine II-3 - painted turtle
C:Species: Chrysemys picta (painted turtle)
C:Date: 08-Nov-1996 #sequence_revision 08-Nov-1996 #text_change 07-May-
C:Accession: D58208

```

06:25:20 2004

us-09-245-198a-4.oligo.rpr

insky, H.E.; Elsev, R.M.; Wright, C.L.; Rice, P.; Bell, J.E.; Sharp, D.
1, 23547-23557, 1996
ies of reptiles.
r: A58208; MUID:96394458; PMID:8798564
08
nary
protein
<HUN>
erm histone
2.5%; Score 7; DB 2; Length 45;
arity 100.0%; Pred. No. 26;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
GRRG 49
|||||
GRRG 30
uinus imperator
us imperator
93 #sequence_revision 12-Apr-1996 #text_change 21-Jul-2000
82
iva, R.
. 19, 5786, 1991
e 1 gene sequence from the primate *Saguinus imperator* isolated with PCR
X: S22582; MUID:92051332; PMID:1840669
82
DNA
<QUE>
s: EMBL:X61678; NID:958405; PIDN:CAA43853.1; PID:94494091
rs translated the codon TAC for residue 43 as Thr
erm histone
osomal protein; DNA binding; nucleus; spermatogenesis
2.5%; Score 7; DB 2; Length 50;
arity 100.0%; Pred. No. 28;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
RGRG 48
|||||
RGRG 24
American opossum
is virginiana; *Didelphis marsupialis virginiana* (North American opossum)
95 #sequence_revision 06-Jan-1995 #text_change 23-Jul-1999
15
Nishikawa, S.; Connor, W.; Dixon, G.H.
215, 63-72, 1993
ization of a marsupial sperm protamine gene and its transcripts from
c: S34045; MUID:93345500; PMID:8344286
15
lary
DNA
<WIN>
s: EMBL:X74044; NID:9407062; PIDN:CAA52193.1; PID:9407063
erm histone
inding; nucleus
2.5%; Score 7; DB 2; Length 58;
arity 100.0%; Pred. No. 32;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
RRG 49
|||||
RRG 41

RESULT 23

F87604
hypothetical protein CC2870 [imported] - *Caulobacter crescentus*
C:Species: *Caulobacter crescentus*
C>Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr-2001
C:Accession: F87604
R:Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen,
B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.
n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter,
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A>Title: Complete Genome Sequence of *Caulobacter crescentus*.
A:Reference number: A87249; MUID:21173698; PMID:11259647
A:Accession: F87604
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-86 <STO>
A:Cross-references: GB:AE005673; NID:gl3424486; PIDN:AAK24834.1; GSFI
C:Genetics:
A:Gene: CC2870

Query Match 2.5%; Score 7; DB 2; Length 86;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 59 LALGLGL 65
|||||
Db 26 LALGLGL 32

RESULT 24

F87993
protein ZC334.3 [imported] - *Caenorhabditis elegans*
C:Species: *Caenorhabditis elegans*
C>Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 10-May-2001
C:Accession: F87993
R:anonymous, The C. elegans Sequencing Consortium.
Science 282, 2012-2018, 1998
A>Title: Genome sequence of the nematode *C. elegans*: a platform for i
A:Reference number: A75000; MUID:99069613; PMID:9851916
A>Note: see websites genome.wustl.edu/gsc/C_elegans/ and www.sanger.a
A>Note: published errata appeared in Science 283, 35, 1999; Science 2
A:Accession: F87993
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-102 <STO>
A:Cross-references: GB:chr I; PIDN:CAB04964.1; PID:g3881432; GSPDB:GN
A>Note: predicted using Genefinder
C:Genetics:
A:Gene: ZC334.3
A:Map position: 1

Query Match 2.5%; Score 7; DB 2; Length 102;
Best Local Similarity 100.0%; Pred. No. 52;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 55 LLVPLAL 61
|||||
Db 33 LLVPLAL 39

RESULT 25

PH1560
IG heavy chain V region (clone VH32) - human (fragment)
C:Species: *Homo sapiens* (man)
C>Date: 05-Aug-1994 #sequence_revision 05-Aug-1994 #text_change 21-Jan-1994
C:Accession: PH1560
R:Rassenti, L.Z.; Kipps, T.J.
J. Exp. Med. 177, 1039-1046, 1993
A>Title: Lack of extensive mutations in the VH5 genes used in common i
A:Reference number: PH1557; MUID:93210459; PMID:7681468
A:Accession: PH1560

```

QA
<RAS>
immunoglobulin V region; immunoglobulin homology
tetramer; immunoglobulin
immunoglobulin homology <IMM>

```

2.5%; Score 7; DB 2; Length 115;
arity 100.0%; Pred. No. 58;
onservative 0; Mismatches 0; Indels

AV 76
|||
AV 11

ical protein PA3557 [imported] - Pseudomonas aeruginosa (strain PA01)
nas aeruginosa
) #sequence revision 15-Sep-2000 #text change 31-Dec-2000

am, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.; Br
: Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lim,
: M.V.
genome sequence of *Pseudomonas aeruginosa* PA01, an opportunistic patho
: AB2950; PMID:20437337; PMID:10984043
1, 2000

 $\text{STO} >$

```

: GB:AE0004776; GB:AE0004091; NID:G9949701; PIDN:AAG06945.1; GSPDB:GN001
: cce: strain PA01

```

```

2.5%; Score 7; DB 2; Length 115;
urity 100.0%; Pred. No. 58;
onservative 0; Mismatches 0; Indels

```

55 GL II 65

```
.n, l2K ~ Escherichia coli insertion sequence IS5
      : hia coli
      : #sequence revision 18-Dec-1981 #text_change 10-Sep-1999
      : : C03582: A04466
```

), M.
.981
)tide sequence of IS5 from *Escherichia coli*.
A91483; MUID:82028653; PMID:6269959

IA
:SCH>
1 Bree.

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IA
:ENG>

insertion sequence IS5
erichia coli hypothetical protein o263

```

2.5%; Score 7; DB 1; Length 118;
arity 100.0%; Pred. No. 59;
conservative 0; Mismatches 0; Indels

```

QY 23 DGGAVRQ 29
Db 107 DGGAVRQ 113

RESULT 28
AE1753

Orf51 [bacteriophage b1285] homolog lin2570 [imported] - *Listeria inn*
C:Species: *Listeria innocua*
C:date: 27-Nov-2001 #nucleotide_revision 27-Nov-2001 #text_change 27-Nov-
C:Accession: AF1753
R:Glaser, P.; Pringle, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Be-
D.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entia
J.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kretz, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.;
ok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.;
A:title: Comparative genomics of *Listeria* species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669

A;Accession: AE1753
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-118 <GLA>

A; Cross-references: GB:AL52022; PIDN:CAC97797.1; PID:gl6415092; GSPDB
A; Experimental source: strain Clipl1262
C; Genetics:
A; Gene: lin2570

```
Query Match      2.5%; Score 7; DB 2; Length 118;
Best Local Similarity 100.0%; Pred. No. 59;
Matches 7; Conservative 0; Mismatches 0; Indels 0; G
```

Qy	2	SLLDFEI	8
Db	61	SLLDFEI	67

RESULT 29
AH2707

conserved hypothetical protein Atul065 [imported] - Agrobacterium tumefaciens
C:Species: Agrobacterium tumefaciens
C:Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 18-Nov
C:Accession: AH2707
R:Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.;
erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutayavin, T.; Levy, R.
; Karp, P.; Romero, P.; Zhang, S.
Science 294, 2317-2323, 2001
A:Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry,
ster, E.W.

A:Title: The Genome of the Natural Genetic Engineer *Agrobacterium tumefaciens* strain C58
A:Reference number: AB2577; MUID:21608550; PMID:11743193
A:Accession: AH2707
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-123 <KUR>
A:Cross-references: GB:AE00688; PIDN:AA42078.1; PID:g17739458; GSPDB:100000000
A:Experimental source: strain C58 (Dupont)
C:Genetics:
A:Gene: Atul065
A:Map position: circular chromosome

Query Match	2.5%;	Score 7;	DB 2;	Length 123;
Best Local Similarity	100.0%;	Pred. No. 61;		
Matches	7;	Conservative	0. Mismatches	0. Indels

Qy	12	RLPLPRS	18
Db	50	RLPLPRS	56

RESULT 30
T27519

127319
hypothetical protein ZC334.3 - *Caenorhabditis elegans*

labditis elegans
#sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
19

EMBL Data Library, November 1996
C: Z20381
A: 19
ary; translated from GB/EMBL/DBJ
DNA
<WIL>
s: EMBL:Z82082; PIDN:CAB04964.2; GSPDB:GN000019; CESP:ZC334.3
rce: clone ZC334
1.3

2.5%; Score 7; DB 2; Length 125;
arity 100.0%; Pred. No. 62;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AL 61
|||
AL 39

at
norvegicus (Norway rat)
#sequence_revision 02-Aug-1996 #text_change 16-Jul-1999
10
uet, E.N.
Res. Commun. 197, 612-618, 1993
of rat interleukin-13 (IL-13) cDNA and analysis of IL-13 gene expression
; 152290; MUID:94092138; PMID:7916615
10
ary; translated from GB/EMBL/DBJ
RNA
<RES>
: GB:L26913; NID:9438875; PIDN:AAA16478.1; PID:9438876

erleukin-13
2.5%; Score 7; DB 2; Length 131;
arity 100.0%; Pred. No. 64;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

CLG 71
|||
CLG 15

protein P600 precursor - mouse
culus (house mouse)
9 #sequence_revision 28-Aug-1989 #text_change 16-Jul-1999
2
awski, S.M.; Mosmann, T.R.; Zurawski, G.
79-687, 1989
of small inducible proteins secreted by leukocytes are members of a ne
ation processes.
: A30552; MUID:89093958; PMID:2521353
2
RNA
<BRO>
: GB:M23504; NID:9533246; PIDN:AAA40149.1; PID:9533247
erleukin-13
2.5%; Score 7; DB 2; Length 131;
arity 100.0%; Pred. No. 64;

Matches 7; Conservative 0; Mismatches 0; Indels 0;
QY 65 LALACLG 71
|||
Db 9 LALACLG 15

RESULT 33
T37116
probable transposase, truncated [imported] - Streptomyces coelicolor
C:Species: Streptomyces coelicolor
C>Date: 08-Sep-2000 #sequence_revision 08-Sep-2000 #text_change 15-Sep
C:Accession: T37116
R:Saunders, D.C.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B
submitted to the EMBL Data Library, August 1999
A:Reference number: Z21588
A:Accession: T37116
A>Status: Preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-146 <SAU>
A:Cross-references: EMBL:AL109950; PIDN:CAB52967.1; GSPDB:GN000070; SC
A:Experimental source: strain A3(2)
C:Genetics:
A:Gene: SCORDB:SCJ4.33c
C:Superfamily: Synchocystis transposase sl11710

Query Match 2.5%; Score 7; DB 2; Length 146;
Best Local Similarity 100.0%; Pred. No. 70;
Matches 7; Conservative 0; Mismatches 0; Indels 0;
QY 70 LGLLIAV 76
|||
Db 102 LGLLIAV 108

RESULT 34
A71217
hypothetical protein PH2001 - Pyrococcus horikoshii
C:Species: Pyrococcus horikoshii
C>Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 20-Jun
C:Accession: A71217
R:Kawarabayashi, Y.; Sawada, M.; Horikawa, H.; Haikawa, Y.; Hino, Y.;
M.; Ohfuku, Y.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.;
DNA Res. 5, 55-76, 1998
A:Title: Complete sequence and gene organization of the genome of a hy
A:Reference number: A71000; MUID:98344137; PMID:9679194
A:Accession: A71217
A>Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-147 <KAW>
A:Cross-references: GB:AP000007; GB:AP000001; NID:g3236134; NID:g32361
A:Experimental source: strain OT3
A:Note: this accession replaces an interim accession for a sequence re
A:Note: this sequence is split into two separate translations in GenBa
C:Genetics:
A:Gene: PH2001
C:Superfamily: Pyrococcus horikoshii hypothetical protein PH2001

Query Match 2.5%; Score 7; DB 2; Length 147;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 67 LALICGL 73
|||
Db 45 LALICGL 51

RESULT 35
T08734
hypothetical protein DKEP566F0546.1 - human (fragment)
C:Species: Homo sapiens (man)
C>Date: 11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 13-Aug
C:Accession: T08734

```

; Obermaier, B.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
rotein Sequence Database, May 1999
4
RNA
<OTT>
: EMBL:AL050075
ce: fetal kidney; clone DKFZp566F0546
546.1
2.5%; Score 7; DB 2; Length 150;
arity 100.0%; Pred. No. 72;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
LGP 240
|||
LGP 38
- rice
ativa (rice)
} #sequence_revision 30-Sep-1993 #text_change 20-Jun-2000
l, H.; Yamada, T.; Tanaka, K.; Takeuchi, S.; Nakamura, R.; Matsuda, T.
l, 239-248, 1993
ture and expression of rice seed allergenic proteins belonging to the
: S31078; MUID:93144699; PMID:7678765
RNA
:ADA>
EMBL:D11430; NID:g218196; PIDN:BAA01996.1; PID:g218197
t alpha-amylase inhibitor
2.5%; Score 7; DB 2; Length 157;
arity 100.0%; Pred. No. 75;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
VS 78
|||
VS 17
tiva (rice)
#sequence_revision 24-Mar-1999 #text_change 16-Jul-1999
J.H.; Eun, M.Y.
BL Data Library, January 1998
otide sequence of rice allergenic protein.
Z14691
IV; translated from GB/EMBL/DBJ
NA
YUN>
EMBL:AF042200; NID:g2827315; PIDN:AAB99797.1; PID:g2827316
ce: strain Nipponbare
t alpha-amylase inhibitor
2.5%; Score 7; DB 2; Length 157;
arity 100.0%; Pred. No. 75;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
VS 78
|||
VS 17

```

```

RESULT 38
A75567
conserved hypothetical protein - Deinococcus radiodurans (strain R1)
C:Species: Deinococcus radiodurans
C>Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 17-Mar
C:Accession: A75567
R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.
Science 286, 1571-1577, 1999
A>Title: Genome sequence of the radioresistant bacterium Deinococcus r
A:Reference number: A75250; MUID:20036896; PMID:10567266
A:Accession: A75567
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-157 <WHI>
A:Cross-references: GB:AE001867; GB:AE000513; NID:g6457693; PIDN:AAF09
A:Experimental source: strain R1
C:Genetics:
A:Gene: DR0033
A:Map position: 1

```

```

Query Match 2.5%; Score 7; DB 2; Length 157;
Best Local Similarity 100.0%; Pred. No. 75;
Matches 7; Conservative 0; Mismatches 0; Indels 0; G
Qy 261 IRTLPA 267
Db 22 IRTLPA 28

```

```

RESULT 39
E75530
hypothetical protein - Deinococcus radiodurans (strain R1)
C:Species: Deinococcus radiodurans
C>Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 31-Mar
C:Accession: E75530
R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.
Science 286, 1571-1577, 1999
A>Title: Genome sequence of the radioresistant bacterium Deinococcus r
A:Reference number: A75250; MUID:20036896; PMID:10567266
A:Accession: E75530
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-157 <WHI>
A:Cross-references: GB:AE001895; GB:AE000513; NID:g6458024; PIDN:AAF09
A:Experimental source: strain R1
C:Genetics:
A:Gene: DR0352
A:Map position: 1

```

```

Query Match 2.5%; Score 7; DB 2; Length 157;
Best Local Similarity 100.0%; Pred. No. 75;
Matches 7; Conservative 0; Mismatches 0; Indels 0; G
Qy 43 RRRGRG 49
Db 144 RRRGRG 150

```

```

RESULT 40
S59925
allergen RASB precursor - rice
C:Species: Oryza sativa (rice)
C>Date: 15-Feb-1996 #sequence_revision 01-Mar-1996 #text_change 20-Jun-
C:Accession: S59925
R:Alvarez, A.M.; Adachi, T.; Nakase, M.; Aoki, N.; Nakamura, R.; Matsuc
Biochim. Biophys. Acta 1251, 201-204, 1995
A>Title: Classification of rice allergenic protein cDNAs belonging to t
A:Reference number: S59922; MUID:95399441; PMID:7669811
A:Accession: S59925

```

```

ary
RNA
<ALV>
: EMBL:D42142; NID:gl398917; PIDN:BAA07713.1; PID:gl398918
at alpha-amylase inhibitor
      2.5%; Score 7; DB 2; Length 160;
larity 100.0%; Pred.No. 76;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AVVS 78
|||||
AVVS 17

rotein hlpA precursor - Escherichia coli (strain K-12)
: DNA-binding 17K protein; histone-like protein hlp
ichia coli
19 #sequence_revision 30-Jun-1989 #text_change 01-Mar-2002
4; A38063; S13728; B64742; I54944; S20426
e, K.
1988
and sequencing of the gene for the DNA-binding 17K protein of Escherich
: J10304; MUID:88329735; PMID:2843433
14
DNA
<HOL>
: GB:M21118; NID:gl47821; PIDN:AAA24630.1; PID:gl47822
13
rotein
<HO2>
rce: strain B
ethaxam, S.
334-344, 1991
and nucleotide sequence of the fira gene and the fira200(Ts) allele fro
: S13728; MUID:91100302; PMID:1987124
18
ary
DNA
- <DIC>
: EMBL:X54797; NID:941468; PIDN:CAA38567.1; PID:g41469
rce: strain K-12, substrain MG1655
Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; Co
lau, B.; Shao, Y.
1462, 1997
ete genome sequence of Escherichia coli K-12.
: A64720; MUID:97426617; PMID:9278503
2
acid sequence not shown; translation not shown
NA
<BLAT>
: GB:AE000127; GB:U00096; NID:gl786370; PIDN:AACT3289.1; PID:gl786375;
rce: strain K-12, substrain MG1655
i, P.; Vaara, M.
1223-1229, 1991
Gene of Yersinia enterocolitica: cloning, sequencing, expression, and
: I54944; MUID:91123198; PMID:1991717
4
ed from GB/EMBL/DBJ
NA
L', 16-148, E', 150-152, I', 154-161 <RES>
: EMBL:X75465; NID:9432661; PIDN:CAA53207.1; PID:g432662
A protein has been believed to be a histone-like constituent of bacter
1-binding 17K protein
ine protein
nal sequence #status predicted <SIG>
outer membrane protein hlpA #status predicted <WAT>
      2.5%; Score 7; DB 1; Length 161;

```

```

Best Local Similarity 100.0%; Pred.No. 77;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY      62 GLGLALA 68
DB      9 GLGLALA 15

RESULT 42
D90651
histone-like protein hlpA [imported] - Escherichia coli (strain O157:)
C:Species: Escherichia coli
C>Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 03-Aug-2001
C:Accession: D90651
R:Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yoko-
gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shina-
DNA Res. 8, 11-22, 2001
A:Title: Complete genome sequence of enterohemorrhagic Escherichia co
A:Reference number: A99629; MUID:21156231; PMID:11258796
A:Accession: D90651
A:Molecule type: DNA
A:Residues: 1-161 <HAV>
A:Cross-references: GB:BA000007; PIDN:BA33603.1; PID:gl3359636; GSPDI
A:Experimental source: strain O157:H7, substrain RIMD 0509952
C:Genetics:
A:Gene: EC80180
C:Superfamily: DNA-binding 17K protein

Query Match      2.5%; Score 7; DB 2; Length 161;
Best Local Similarity 100.0%; Pred.No. 77;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY      62 GLGLALA 68
DB      9 GLGLALA 15

RESULT 43
D85502
hypothetical protein hlpA [imported] - Escherichia coli (strain O157:)
C:Species: Escherichia coli
C>Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001
C:Accession: D85502
R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.;
Miller, L.; Grobeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potam-
Nature 409, 529-533, 2001
A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H
A:Reference number: A85480; MUID:21074935; PMID:11206551
A:Accession: D85502
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-161 <STO>
A:Cross-references: GB:AB005174; NID:gl2512906; PIDN:AAG54480.1; GSPDI
A:Experimental source: strain O157:H7, substrain EDD933
C:Genetics:
A:Gene: hlpA
C:Superfamily: DNA-binding 17K protein

Query Match      2.5%; Score 7; DB 2; Length 161;
Best Local Similarity 100.0%; Pred.No. 77;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY      62 GLGLALA 68
DB      9 GLGLALA 15

RESULT 44
T24937
hypothetical protein W03C9.4 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 29-Oct-1999

```

```

7; T26123
WBL Data Library, July 1995
7
A; Accession: S76920
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-164 <KAN>
A; Note: the nucleotide sequence was submitted to the EMBL Data Library
Query Match 2.5%; Score 7; DB 2; Length 164;
Best Local Similarity 100.0%; Pred. No. 78;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 104 SELNPQT 110
Db 37 SELNPQT 43
RESULT 47
S44789
D2007.4 protein - Caenorhabditis elegans
C; Species: Caenorhabditis elegans
C; Date: 14-Sep-1994 #sequence_revision 12-May-1995 #text_change 23-Mar
C; Accession: S44789
R; Favell, A.D.
submitted to the EMBL Data Library, May 1993
A; Description: Sequence of the C. elegans cosmid D2007.
A; Reference number: S44619
A; Accession: S44789
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-170 <FAV>
A; Cross-references: EMBL:L16560; NID:g289666; PID:g289670
C; Genetics:
A; Introns: 43/2; 121/3
Query Match 2.5%; Score 7; DB 2; Length 170;
Best Local Similarity 100.0%; Pred. No. 80;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 221 VLALRCL 227
Db 94 VLALRCL 100
RESULT 48
D87638
transcription regulator, GntR family [imported] - Caulobacter crescentus
C; Species: Caulobacter crescentus
C; Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr
C; Accession: D87638
R; Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen,
B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.;
n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter,
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A; Title: Complete Genome Sequence of Caulobacter crescentus.
A; Reference number: A87249; MUID:21173698; PMID:11259647
A; Accession: D87638
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-174 <STO>
A; Cross-references: GB:AE005673; NID:gl3424808; PIDN:AAK25104.1; GSPDB
C; Genetics:
A; Gene: CC3142
Query Match 2.5%; Score 7; DB 2; Length 174;
Best Local Similarity 100.0%; Pred. No. 82;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 138 ARRATAA 144
Db 151 ARRATAA 157
7; T26123
WBL Data Library, July 1995
7
A; Accession: S76920
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-164 <KAN>
A; Note: the nucleotide sequence was submitted to the EMBL Data Library
Query Match 2.5%; Score 7; DB 2; Length 164;
Best Local Similarity 100.0%; Pred. No. 78;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 104 SELNPQT 110
Db 37 SELNPQT 43
RESULT 47
S44789
D2007.4 protein - Caenorhabditis elegans
C; Species: Caenorhabditis elegans
C; Date: 14-Sep-1994 #sequence_revision 12-May-1995 #text_change 23-Mar
C; Accession: S44789
R; Favell, A.D.
submitted to the EMBL Data Library, May 1993
A; Description: Sequence of the C. elegans cosmid D2007.
A; Reference number: S44619
A; Accession: S44789
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-170 <FAV>
A; Cross-references: EMBL:L16560; NID:g289666; PID:g289670
C; Genetics:
A; Introns: 43/2; 121/3
Query Match 2.5%; Score 7; DB 2; Length 170;
Best Local Similarity 100.0%; Pred. No. 80;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 221 VLALRCL 227
Db 94 VLALRCL 100
RESULT 48
D87638
transcription regulator, GntR family [imported] - Caulobacter crescentus
C; Species: Caulobacter crescentus
C; Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr
C; Accession: D87638
R; Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen,
B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.;
n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter,
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A; Title: Complete Genome Sequence of Caulobacter crescentus.
A; Reference number: A87249; MUID:21173698; PMID:11259647
A; Accession: D87638
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-174 <STO>
A; Cross-references: GB:AE005673; NID:gl3424808; PIDN:AAK25104.1; GSPDB
C; Genetics:
A; Gene: CC3142
Query Match 2.5%; Score 7; DB 2; Length 174;
Best Local Similarity 100.0%; Pred. No. 82;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 138 ARRATAA 144
Db 151 ARRATAA 157

```

n precursor - goat
aegagrus hircus (domestic goat)
85 #sequence revision 12-Apr-1996 #text_change 22-Jun-1999
20; S14507; S42800; S42801
unitzer, G.; Schrank, B.; Stangl, A.
Physiol. Chem. 360, 1595-1604, 1979
o acid sequence of goat beta-lactoglobulin.
A: A91682; MUID:80070611; PMID:511095
20
protein
0 <PRE>
11, A.; Sanchez, A.
EMBL Data Library, March 1991
r: S14507
07
mRNA
<POL>
s: EMBL:X58471; NID:g967; PIDN:CAA41385.1; PID:g968
EMBL Data Library, January 1993
r: S42800
00
mRNA
<KIM>
s: EMBL:Z19569; NID:g437751; PIDN:CAA79623.1; PID:g437752
01
mRNA
<K12>
s: EMBL:Z19570; NID:g437753; PIDN:CAA79624.1; PID:g437754
physiological conditions beta-lactoglobulin exists as an equilibrium mi
socalin; lipocalin homology
gnal sequence #status predicted <SIG>
beta-lactoglobulin #status predicted <MAT>
lipocalin homology <LIP>
Disulfide bonds: #status predicted
2.5%; Score 7; DB 1; Length 180;
arity 100.0%; Pred. No. 84;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ALAC 69
|||||
ALAC 14
1 precursor - sheep
beta-lactoglobulin A; beta-lactoglobulin B; beta-lactoglobulin C; bet
entalis aries, Ovis ammon aries (domestic sheep)
35 #sequence revision 19-Apr-1996 #text_change 22-Jun-1999
19; JQ0748; A30011; B30011; S02136; A03221; S04955
ghan, M.; Simons, J.P.; Clark, A.J.
1990
sation of the alleles encoding ovine beta-lactoglobulins A and B.
s: JQ0748; MUID:91007276; PMID:1976573
19
DNA
<ALIB>
s: GB:M32232
rce: beta-lactoglobulin B
18
DNA
'Y', 39-180 <ALIA>
s: GB:M32232
rce: beta-lactoglobulin A
A.J.
415-426, 1988
ization of the gene encoding ovine beta-lactoglobulin. Similarity to d

A:Reference number: A92942; MUID:88172489; PMID:3351935
A:Accession: A30011
A:Molecule type: DNA
A:Residues: 1-180 <ALII>
A:Cross-references: GB:X14971
A:Experimental source: beta-lactoglobulin I
A:Accession: B30011
A:Molecule type: DNA
A:Residues: 1-37, 'Y', 39-102, 'N', 104-180 <ALII2>
A:Cross-references: GB:X07009
A:Experimental source: beta-lactoglobulin II
R; Harris, S.; Ali, S.; Anderson, S.; Archibald, A.L.; Clark, A.J.
Nucleic Acids Res. 16, 10379-10380, 1988
A:Title: Complete nucleotide sequence of the genomic ovine beta-lacto
A:Reference number: S02136; MUID:89057492; PMID:3194215
A:Accession: S02136
A:Status: translation not shown
A:Molecule type: DNA
A:Residues: 1-180 <HAR>
A:Cross-references: EMBL:X12817; NID:gl313; PIDN:CAA31305.1; PID:gl31
R; Gaye, P.; Hue-Delahaie, D.; Mercier, J.C.; Soulier, S.; Vilotte, J.
Biochimie 68, 1097-1107, 1986
A:Title: Ovine beta-lactoglobulin messenger RNA: nucleotide sequence
A:Reference number: A25136; MUID:87049827; PMID:3096387
A:Accession: A25136
A:Molecule type: mRNA
A:Residues: 1-180 <GAY>
A:Cross-references: GB:X04520; NID:gl315; PIDN:CAA28204.1; PID:gl316
R; Preaux, G.; Braunitzer, G.; Kolde, H.J.
Arch. Int. Physiol. Biochim. 88, B45-B46, 1980
A:Title: Primary structure of ovine beta-lactoglobulin.
A:Reference number: A03221; MUID:80219294; PMID:6155855
A:Accession: A03221
A:Molecule type: protein
A:Residues: 19-37, 'Y', 39-180 <PRE>
R; Brhardt, G.; Godovac-Zimmermann, J.; Conti, A.
Biol. Chem. Hoppe-Seyler 370, 757-762, 1989
A:Title: Isolation and complete primary sequence of a new ovine wild-
A:Reference number: S04955; MUID:89374823; PMID:2775495
A:Accession: S04955
A:Molecule type: protein
A:Residues: 19-37, 'Y', 39-165, 'Q', 167-180 <ERH>
A:Experimental source: beta-lactoglobulin C
C:Comment: This protein is the major milk whey protein of ruminants a
C:Genetics: Under physiological conditions beta-lactoglobulin exists a
A:Gene: BLG
A:Introns: 32/3; 79/2; 104/1; 141/1; 176/1
C:Superfamily: lipocalin; lipocalin homology
C:Keywords: milk; polymorphism
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-180/Product: beta-lactoglobulin #status experimental <MAT>
F:28-178/Domain: lipocalin homology <LIP>
F:84-178, 124-137/Disulfide bonds: #status predicted
Query Match 2.5%; Score 7; DB 1; Length 180;
Best Local Similarity 100.0%; Pred. No. 84;
Matches 7; Conservative 0; Mismatches 0; Indels 0;
QY 63 LGLALAC 69
DB 8 LGLALAC 14
RESULT 51
B60738
insulin-like growth factor II precursor - pig
C:Species: Sus scrofa domestica (domestic pig)
C>Date: 28-Apr-1993 #sequence_revision 30-Sep-1993 #text_change 13-Nov
C:Accession: S12614; B60738
R; Catchpole, I.R.; Engstroem, W.
Nucleic Acids Res. 18, 6430, 1990
A:Title: Nucleotide sequence of a porcine insulin-like growth factor I

C:Genetics:

A:Map position: 3

A:Introns: 42/2; 98/3; 126/3; 176/3

A:Note: C35D10.8

C:Superfamily: Caenorhabditis elegans hypothetical protein C35D10.8

Query Match 2.5%; Score 7; DB 2; Length 206;

Best Local Similarity 100.0%; Pred. No. 94;

Matches 7; Conservative 0; Mismatches 0; Indels 0; G 0;

QY 136 TRARRAI 142

DB 98 TRARRAI 104

|||||

RESULT 54

AH0692

conserved hypothetical protein ydgQ [imported] - Salmonella enterica s

C:Species: Salmonella enterica subsp. enterica serovar Typhi

A:Note: this species has also been called Salmonella typhi

C:Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 18-Nov

C:Accession: AH0692

R:Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; W

th, T.; Connerton, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.;

S.; Moule, S.; O'Gaora, P.

Nature 413, 848-852, 2001

A:Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton

A:Title: Complete genome sequence of a multiple drug resistant *Salmonella*

A:Reference number: AB0502; MUID:21534947; PMID:11677608

A:Accession: AH0692

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-230 <PAR>

A:Cross-references: GB:AL513382; PIDN:CAD01913.1; PID:gl6502755; GSPD8

C:Genetics:

A:Gene: ydgQ

C:Superfamily: conserved hypothetical protein H11688

Query Match 2.5%; Score 7; DB 2; Length 230;

Best Local Similarity 100.0%; Pred. No. 1e+02;

Matches 7; Conservative 0; Mismatches 0; Indels 0; G 0;

QY 60 ALGLGLA 66

DB 38 ALGLGLA 44

|||||

RESULT 55

B64920

probable membrane protein ydgQ - Escherichia coli (strain K-12)

C:Species: Escherichia coli

C:Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 01-Mar

C:Accession: B64920

R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland,

A.; Rose, D.J.; Mau, B.; Shao, Y.

Science 277, 1453-1462, 1997

A:Title: The complete genome sequence of *Escherichia coli* K-12.

A:Reference number: A64720; MUID:97426617; PMID:9278503

A:Accession: B64920

A>Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-231 <BLAT>

A:Cross-references: GB:AE000258; GB:U00096; NID:g2367121; PIDN:AACT4704

A:Experimental source: strain K-12, substrain MG1655

C:Genetics:

A:Gene: ydgQ

C:Superfamily: conserved hypothetical protein H11688

C:Keywords: transmembrane protein

C:Keywords: transmembrane #status predicted <TM01>

C:Keywords: transmembrane #status predicted <TM02>

C:Keywords: transmembrane #status predicted <TM03>

C:Keywords: transmembrane #status predicted <TM04>

```

transmembrane #status predicted <TM05>
transmembrane #status predicted <TM06>

      2.5%; Score 7; DB 2; Length 231;
Identity 100.0%; Pred. No. 1e+02;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LGLA 66
|||||
LGLA 44

ein ECs2341 [imported] - Escherichia coli (strain O157:H7, substrain R1)
ichia coli
01 #sequence_revision 18-Jul-2001 #text_change 03-Aug-2001
21
kino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.;
unaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
, 2001
genome sequence of enterohemorrhagic Escherichia coli O157:H7 and gen
r: A99629; MUID:21156231; PMID:11258796
21
nary
DNA
<HAY>
s: GB:BA000007; PIDN:BA035764.1; PID:gl3361808; GSPDB:GN00154
urce: strain O157:H7, substrain RMD 050952

served hypothetical protein HI1688

      2.5%; Score 7; DB 2; Length 231;
Identity 100.0%; Pred. No. 1e+02;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LGLA 66
|||||
LGLA 44

sin ydgQ [imported] - Escherichia coli (strain O157:H7, substrain EDL93)
ichia coli
01 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001
70
unkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew
rk, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
13, 2001
sequence of enterohemorrhagic Escherichia coli O157:H7.
r: AB5480; MUID:21074935; PMID:11206551
70
nary
DNA
<STO>
r: GB:AE005174; NID:gl2515621; PIDN:AAG56621.1; GSPDB:GN00145; UWGP:Z26
urce: strain O157:H7, substrain EDL933

served hypothetical protein HI1688

      2.5%; Score 7; DB 2; Length 231;
Identity 100.0%; Pred. No. 1e+02;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LGLA 66
|||||
LGLA 44

```

```

C86665
amino acid ABC transporter permease protein [imported] - Lactococcus l
C:Species: Lactococcus lactis subsp. lactis
C:Date: 23-Mar-2001 #sequence_revision 23-Mar-2001 #text_change 09-Dec
C:Accession: C86665
R;Bolotin, A.; Wincker, P.; Mauger, S.; Jalllon, O.; Malarne, K.; Wei
Genome Res. 11, 731-753, 2001
A;Title: The complete genome sequence of the lactic acid bacterium La
A;Reference number: A86625; MUID:21235186; PMID:11337471
A;Accession: C86665
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-231 <STO>
A;Cross-references: GB:AE005176; PID:gl2723189; PIDN:AAK04421.1; GSPD
A;Experimental source: strain IL1403
C:Genetics:
A;Gene: ydCC
C;Superfamily: ABC transporter permease protein

Query Match      2.5%; Score 7; DB 2; Length 231;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY      56 LVPLALG 62
DB      107 LVPLALG 113

RESULT 59
S60767
ribonuclease III - Coxiella burnetii
C:Species: Coxiella burnetii
C:Date: 27-Apr-1996 #sequence_revision 13-Mar-1997 #text_change 22-Jun
C:Accession: S60767
R;Zuber, M.; Hoover, T.A.; Powell, B.S.; Court, D.L.
Mol. Microbiol. 14, 291-300, 1994
A;Title: Analysis of the rnc locus of Coxiella burnetii.
A;Reference number: S60767; MUID:95131751; PMID:7830573
A;Accession: S60767
A;Status: preliminary; nucleic acid sequence not shown
A;Molecule type: DNA
A;Residues: 1-233 <ZUB>
A;Cross-references: EMBL:L27436; NID:g439870; PIDN:AAA69690.1; PID:g4
C;Superfamily: ribonuclease III; double-stranded RNA-binding repeat h
F,150-223/Domain: double-stranded RNA-binding repeat homology <DSR>

Query Match      2.5%; Score 7; DB 2; Length 233;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY      10 ARRLPLP 16
DB      164 ARRLPLP 170

RESULT 60
AB0273
probable membrane protein YPO2240 [imported] - Yersinia pestis (strain
C:Species: Yersinia pestis
C:Date: 02-Nov-2001 #sequence_revision 02-Nov-2001 #text_change 09-Nov
C:Accession: AB0273
R;Parkhill, J.; Wren, B.W.; Thomson, N.R.; Titball, R.W.; Holden, M.T.
deno-Tarraga, A.M.; Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis
il, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; White
Nature 413, 523-527, 2001
A;Title: Genome sequence of Yersinia pestis, the causative agent of p
A;Reference number: AB00001; MUID:21470413; PMID:11586360
A;Accession: AB0273
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-233 <KUR>
A;Cross-references: GB:AL590842; PIDN:CAC91046.1; PID:gl5980240; GSPDE
C:Genetics:

```

erved hypothetical protein H11688
 2.5%; Score 7; DB 2; Length 233;
 rity 100.0%; Pred. No. 1.1e+02;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 LA 66
 ||
 LA 44
 synthase-like protein [imported] - Arabidopsis thaliana
 sis thaliana (mouse-ear cress)
 #sequence_revision 16-Feb-2001 #text_change 17-May-2002
 ropan Union Arabidopsis Genome Sequencing Consortium, The Cold Spring
 , 1999
 nd analysis of chromosome 4 of the plant Arabidopsis thaliana.
 A85001; MUID:20083488; PMID:10617198
 ry
 A
 STO>
 GB_NC_001268; NID:g7267660; PIDN:CAB78088.1; GSPDB:GN00140
 ransporting ATP synthase delta chain
 2.5%; Score 7; DB 2; Length 234;
 rity 100.0%; Pred. No. 1.1e+02;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 SS 237
 ||
 SS 54
 nslocating NADH dehydrogenase (ubiquinone) (EC 1.6.5.-) nqrD chain H1
 lus influenzae
 #sequence_revision 18-Aug-1995 #text_change 21-Jul-2000
 Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage, A
 ott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman, J
 C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghegan, N.S.M.
 2, 1995
 .L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter,
 me random sequencing and assembly of Haemophilus influenzae Rd.
 A64000; MUID:95350630; PMID:7542800
 cid sequence not shown; translation not shown
 IA
 TIGR>
 GB:U32841; GB:L42023; NID:gl574529; PIDN:AAC23334.1; PID:gl574540; T
 erved hypothetical protein H11688
 ductase
 2.5%; Score 7; DB 2; Length 235;
 rity 100.0%; Pred. No. 1.1e+02;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 ILA 66
 ||
 ILA 68

probable cytochrome b-561 [imported] - Arabidopsis thaliana
 C:Species: Arabidopsis thaliana (mouse-ear cress)
 C:Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-
 C:Accession: A86387
 R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; W
 Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy,
 ansen, N.F.; Hughes, B.; Huizar, L.
 Nature 408, 816-820, 2000
 A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Kha
 A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Mai
 Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
 A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.;
 ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
 A>Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis
 A:Reference number: A86141; MUID:21016719; PMID:11130712
 A:Accession: A86387
 A>Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-236 <STO>
 A:Cross-references: GB:A8005172; NID:gil079498; PIDN:AAG29209.1; GSPDB:
 C:Genetics:
 A:Map position: 1

Query Match 2.5%; Score 7; DB 2; Length 236;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps
 QY 61 LGLGLAL 67
 Db 194 LGLGLAL 200

RESULT 64
 H70866
 hypothetical protein Rv2473 - Mycobacterium tuberculosis (strain H37RV)
 C:Species: Mycobacterium tuberculosis
 C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-
 C:Accession: H70866
 R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Har
 Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squ
 Nature 393, 537-544, 1998
 A:Authors: Squires, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell
 A>Title: Deciphering the biology of Mycobacterium tuberculosis from the
 A:Reference number: A70500; MUID:98295987; PMID:96344230
 A:Accession: H70866
 A>Status: preliminary; nucleic acid sequence not shown; translation not
 A:Molecule type: DNA
 A:Residues: 1-238 <COL>
 A:Cross-references: GB:AL021246; GB:AL123456; NID:g3261507; PIDN:CAA161
 A:Experimental source: strain H37RV
 C:Genetics:
 A:Gene: RV2473

Query Match 2.5%; Score 7; DB 2; Length 238;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps
 QY 74 LAVVSLG 80
 Db 90 LAVVSLG 96

RESULT 65
 G83208
 conserved hypothetical protein PA3494 [imported] - Pseudomonas aerugin
 C:Species: Pseudomonas aeruginosa
 C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 27-Nov-
 C:Accession: G83208
 R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.
 adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.
 ; Lory, S.; Olson, M.V.
 Nature 406, 959-964, 2000

genome sequence of *Pseudomonas aeruginosa* PA01, an opportunistic pathogen
 : AB2950; MUID:2043737; PMID:10984043

Query Match 2.5%; Score 7; DB 2; Length 240;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 0; Mismatches 0; Indels 0; Gaps 0;
 : ST0>
 : GB:AE004770; GB:AE004091; NID:99949633; PIDN:AA06882.1; GSPDB:GN001
 cce: strain PA01

served hypothetical protein H1688

Query Match 2.5%; Score 7; DB 2; Length 240;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 0; Mismatches 0; Indels 0; Gaps 0;

GLA 66

GLA 46

synthesis protein moeB - Haemophilus influenzae (strain Rd KW20)

5 #sequence_revision 18-Aug-1995 #text_change 29-Sep-1999

: Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage, A.
 :; Cott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman, J.
 :; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghagen, N.S.M.
 : 12, 1995

: L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter,
 :ome random sequencing and assembly of Haemophilus influenzae Rd.
 : : A64000; MUID:95350630; PMID:7542800

acid sequence not shown; translation not shown

NA

: GB:U032823; GB:142023; NID:G1574281; PIDN:AAC23099.1; PID:G1574288; T

pterin biosynthesis
 ydopterin biosynthesis protein moeB
 ienum; molybdopterin biosynthesis

Query Match 2.5%; Score 7; DB 2; Length 243;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 0; Mismatches 0; Indels 0; Gaps 0;

NRQ1 185

NRQ1 16

ein ZC196.8 - *Caenorhabditis elegans*
 habditis elegans

99 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999

EMBL Data Library, April 1997
 e sequence of *C. elegans* cosmid ZC196.

r: Z20115

nary; translated from GB/EMBL/DBDJ

DNA

:s: EMBL:U97007; PIDN:AA052298.1; GSPDB:GN00023; CESP:ZC196.8

source: strain Bristol N2; clone ZC196

6.8

A;Map position: 5
 A;Introns: 41/1; 84/2; 144/3; 195/3

Query Match 2.5%; Score 7; DB 2; Length 243;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G

Qy 234 AASSLGP 240

Db 225 AASSLGP 231

RESULT 68

A46066

lymphotoxin beta - human

C;Species: Homo sapiens (man)

C;Date: 21-Sep-1993 #sequence_revision 18-Nov-1994 #text_change 21-Jul

C;Accession: A46066

R;Browning, J.L.; Ngam-ek, A.; Lawton, P.; DeMarinis, J.; Tizard, R.;

Cell 72, 847-856, 1993

A;Title: Lymphotoxin beta, a novel member of the TNF family that forms

A;Reference number: A46066; MUID:93208881; PMID:7916655

A;Accession: A46066

A;Status: preliminary

A;Molecule type: DNA; protein

A;Residues: 1-244 <BRO>

A;Cross-references: GB:111015; NID:G922276; PIDN:AAA36191.1; PID:G2922

A;Note: sequence extracted from NCBI backbone (NCBIN:128086, NCBI:128

C;Keywords: transmembrane protein

Query Match 2.5%; Score 7; DB 2; Length 244;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;

Matches 7; Conservative 0; Mismatches 0; Indels 0; C

Qy 194 GLYLYLC 200

Db 131 GLYLYLC 137

RESULT 69

B32352

molybdopterin biosynthesis protein moeB - *Escherichia coli* (strain K-

N;Alternate names: molybdopterin-converting factor chlN

C;Species: *Escherichia coli*

C;Date: 07-Jun-1990 #sequence_revision 07-Jun-1990 #text_change 01-Ma

C;Accession: B32352; B64820

R;Nohno, T.; Kasai, Y.; Saito, T.

J. Bacteriol. 170, 4097-4102, 1988

A;Title: Cloning and sequencing of the *Escherichia coli* chlN operon

A;Reference number: A32352; MUID:88314906; PMID:3045084

A;Accession: B32352

A;Status: not compared with conceptual translation

A;Molecule type: DNA

A;Residues: 1-249 <NOH>

A;Cross-references: GB:M21151; NID:G145538; PIDN:AAA23580.1; PID:G145

R;Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland

.A.; Rose, D.J.; Mau, B.; Shao, Y.

Science 277, 1453-1462, 1997

A;Title: The complete genome sequence of *Escherichia coli* K-12.

A;Reference number: A64720; MUID:97426617; PMID:9278503

A;Accession: B64820

A;Status: preliminary; nucleic acid sequence not shown; translation n

A;Molecule type: DNA

A;Residues: 1-249 <BLAT>

A;Cross-references: GB:AE000185; GB:U00096; NID:G1787047; PIDN:AACT739

A;Experimental source: strain K-12, substrain MG1655

C;Genetics:

A;Gene: moeB; chlN

A;Map position: 18 min

C;Function:

A;Pathway: molybdopterin biosynthesis

C;Superfamily: molybdopterin biosynthesis protein moeB

C;Keywords: molybdenum; molybdopterin biosynthesis

16:25:20 2004

us-09-245-198a-4.oligo.rpr

```

2.5%; Score 7; DB 2; Length 249;
urity 100.0%; Pred. No. 1.1e+02;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

RQI 185
|||
RQI 16

ynthesis MoeB protein [imported] - Escherichia coli (strain O157:H7, s
chia coli
1 #sequence_revision 18-Jul-2001 #text_change 03-Aug-2001
1
ino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.
naga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
2001
genome sequence of enterohemorrhagic Escherichia coli O157:H7 and genc
: A99629; MUID:21156231; PMID:11258796
1
ary
NA
<HAY>
: GB:BA000007; PIDN:BA34327.1; PID:gl3360363; GSPDB:GN00154
rce: strain O157:H7, substrain RMD 0509952

ybdopterin biosynthesis protein moeB
2.5%; Score 7; DB 2; Length 249;
arity 100.0%; Pred. No. 1.1e+02;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

RQI 185
|||
RQI 16

ynthesis [imported] - Escherichia coli (strain O157:H7, substrain EDL9
ichia coli
01 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001
32
nnett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew
ck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
33, 2001
sequence of enterohemorrhagic Escherichia coli O157:H7.
r: A85480; MUID:21074935; PMID:11206551
92
nary
DNA
<STO>
s: GB:AE005174; MID:gl21513829; PIDN:BA55199.1; GSPDB:GN00145; UWGP:Z10
urce: strain O157:H7, substrain EDL933

lybdopterin biosynthesis protein moeB
2.5%; Score 7; DB 2; Length 249;
larity 100.0%; Pred. No. 1.1e+02;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

NRQI 185
|||
NRQI 16

ynthesis MoeB protein [imported] - Salmonella enterica subsp. enterica

```

```

C;Species: Salmonella enterica subsp. enterica serovar Typhi
A;Note: this species has also been called Salmonella typhi
C;Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 18-Nov
C;Accession: AI0602
R;Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; W
th, T.; Connerton, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.;
, S.; Moule, S.; O'Gaora, P.
Nature 413, 848-852, 2001
A;Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton
A;Title: Complete genome sequence of a multiple drug resistant Salmone
A;Reference number: AB0502; MUID:21534947; PMID:11677608
A;Accession: AI0602
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-249 <PAR>
A;Cross-references: GB:AL513382; PIDN:CAD05291.1; PID:gl6502055; GSPDE
C;Genetics:
A;Gene: STY0884
C;Superfamily: molybdopterin biosynthesis protein moeB

Query Match 2.5%; Score 7; DB 2; Length 249;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 179 LRYNRQI 185
|||
DB 10 LRYNRQI 16

RESULT 73
AI3154
IS21 family transposase istB [imported] - Agrobacterium tumefaciens (i
C;Species: Agrobacterium tumefaciens
C;Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 18-No
C;Accession: AT3154
R;Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E
erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutayavin, T.; Levy, J
; Karp, P.; Romero, P.; Zhang, S.
Science 294, 2317-2323, 2001
A;Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry
ster, E.W.
A;Title: The Genome of the Natural Genetic Engineer Agrobacterium tum
A;Reference number: AB2577; MUID:21608550; PMID:11743193
A;Accession: AI3154
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-252 <KUR>
A;Cross-references: GB:AE008689; PIDN:AAL45655.1; PID:gl7743380; GSPD
A;Experimental source: strain C58 (Dupont)
C;Genetics:
A;Gene: istB
A;Map position: linear chromosome
C;Superfamily: DNA replication protein dnaC

Query Match 2.5%; Score 7; DB 2; Length 252;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 60 ALGLGLA 66
|||
DB 120 ALGLGLA 126

RESULT 74
H98132
istB protein (AJ238712) [imported] - Agrobacterium tumefaciens (strai
C;Species: Agrobacterium tumefaciens
C;Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 18-Nc
C;Accession: H98132
R;Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Qi
A.; Liu, F.; Wollam, C.; Allinger, M.; Doughty, D.; Scott, C.; Lapp
Science 294, 2323-2328, 2001
A;Title: Genome Sequence of the Plant Pathogen and Biotechnology Agei

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c: A97359; MUID:21608551; PMID:11743194
32
nary
DNA
<KUR>
s: GB:AE007870; PIDN:AAK88586.1; PID:gl5158297; GSPDB:GN00170

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2.5%; Score 7; DB 2; Length 252;
larity 100.0%; Pred.No.1.1e+02;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LGLA 66
||||
LGLA 126

synthesis MoeB protein VCA0618 [imported] - Vibrio cholerae (strain N16
cholerae
00 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001
39
; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R.J.;
olaeva, M.D.; Vamathevan, J.; Bass, S.; Qin, H.; Dragoi, I.; Sellers, B.
s, J.J.; Venter, J.C.; Fraser, C.M.
B3, 2000
ence of both chromosomes of the cholera pathogen Vibrio cholerae.
r: A82035; MUID:20406833; PMID:10952301
39
nary
DNA
<HEI>
s: GB:AE004391; GB:AE003853; NID:g9658015; PIDN:AAF96519.1; GSPDB:GN001
urce: serogroup O1; strain N16961; biotype El Tor

lydopterin biosynthesis protein moeB
2.5%; Score 7; DB 2; Length 254;
larity 100.0%; Pred.No.1.1e+02;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

NRQI 185
||||
NRQI 18

April 7, 2004, 17:59:25
s

GenCore version 5.1.6
 Copyright (c) 1993 - 2004 Compugen Ltd.
 .n search, using sw model
 .fil 7, 2004, 17:48:13 ; Search time 17 Seconds
 (without alignments)
 869.878 Million cell updates/sec
 -09-245-198a-4
 i
 MSLLDFEISARRLPLRSLG.....PWAHLXAAPLTYFGLFQVH 284

GO
 pop 60.0 , Gapext 60.0
 [681 seqs, 52070155 residues

s satisfying chosen parameters: 141681

h: 0
 fth: 2000000000

sting first 100 summaries

visProt_42:*

the number of results predicted by chance to have a
 r than or equal to the score of the result being printed,
 ad by analysis of the total score distribution.

SUMMARIES

seq	ch	Length	DB	ID	Description
7.7	249	1	TN12	HUMAN	O43508 homo sapien
1.3	225	1	TN12	MOUSE	O54907 mus musculu
3.5	111	1	YFBW	ECOLI	Q47377 escherichia
3.2	733	1	PSAB	ODOSI	P49480 odontella s
3.2	734	1	PSAB	CYACA	Q9t1g6 cyanidium c
2.8	179	1	ADHS	GLUOX	O05544 gluconobact
2.8	220	1	Y304	BRUME	O8vd73 brucella me
2.8	220	1	Y3J1	BRUSU	O8fv59 brucella su
2.8	317	1	MSHR	PANTR	Q9ruk4 pan troglod
2.8	379	1	FDHB	METJA	Q60316 methanococc
2.8	422	1	FP3	MESAU	P23491 mesocricetu
2.8	576	1	CYDC	HAIN	P45081 haemophilus
2.8	586	1	GGT5	HUMAN	P36269 homo sapien
2.5	24	1	HSP3	OCTVU	P83215 octopus vul
2.5	30	1	HSP5	OCTVU	P83217 octopus vul
2.5	49	1	HSP1	SAGIM	P24714 saguinus im
2.5	57	1	HSP1	DIDMA	P35305 didelphis m
2.5	115	1	A62F	DROME	O46202 drosophila
2.5	118	1	Y151	ECOLI	P03838 escherichia
2.5	131	1	IL13	MOUSE	P20109 mus musculu
2.5	131	1	IL13	RAT	P42203 rattus norv
2.5	147	1	YK01	PYRHO	O57781 pyrococcus
2.5	150	1	TNFC	PIG	Q9tsv8 sus scrofa
2.5	157	1	RA05	ORYSA	O01881 oryza sativ
2.5	161	1	HLPA	ECOLI	P11457 escherichia
2.5	170	1	YIM4	CABEL	P34378 caenorhabdi
2.5	180	1	LACB	BUBBU	P02755 bubalus bub
2.5	180	1	LACB	CAPHI	P02756 capra hircu
2.5	181	1	LACB	SHEEP	P02757 ovis aries
2.5	181	1	APT	SHEON	Q8efg1 shewanella
2.5	217	1	DEF1	BIFLO	Q8g534 bifidobacte
2.5	230	1	RNFE	SALTY	Q8xex9 salmonella
2.5	231	1	RNFE	ECOE57	P58344 escherichia

34	7	2.5	231	1	RNFE	ECOLI	P77179 esch
35	7	2.5	233	1	RNC	COXBU	P51837 coxi
36	7	2.5	233	1	RNFE	YERPE	Q8zed4 yers
37	7	2.5	235	1	RNFE	HAEIN	Q57020 haem
38	7	2.5	239	1	TN14	MOUSE	Q9qyh9 mus
39	7	2.5	240	1	RNFE	PSEAE	Q9hyh5 pseu
40	7	2.5	243	1	MOEB	HAEIN	P45211 haem
41	7	2.5	244	1	TNFC	HUMAN	Q06643 homo
42	7	2.5	244	1	TNFC	PANTR	Q86227 pan
43	7	2.5	249	1	LFTF	XANCP	Q8p996 xant
44	7	2.5	249	1	MOEB	ECOLI	P12282 esch
45	7	2.5	249	1	MOEB	SALTY	Q56087 salm
46	7	2.5	257	1	KDIX	SERMA	Q54435 serr
47	7	2.5	277	1	CN09	HUMAN	Q86t03 homo
48	7	2.5	310	1	TNFC	MARMO	Q9jml0 marm
49	7	2.5	316	1	ISPH	XANCP	Q8pbq4 xant
50	7	2.5	316	1	ISPH	XLFA	Q9pas9 xyle
51	7	2.5	324	1	ODPB	BAGSU	P21882 baci
52	7	2.5	329	1	SRA6	CABEL	Q09208 caen
53	7	2.5	335	1	IAG2	RAT	Q35777 ratt
54	7	2.5	344	1	LEU3	THEAQ	P24098 ther
55	7	2.5	352	1	LEU3	DEIRA	Q9rth9 dein
56	7	2.5	357	1	G6PT	MOUSE	P35576 mus
57	7	2.5	357	1	G6PT	RAT	P43428 ratt
58	7	2.5	358	1	PONT	RABIT	P27170 oryc
59	7	2.5	365	1	NQ08	THETH	Q60019 ther
60	7	2.5	394	1	BENE	ACICA	P07775 acin
61	7	2.5	396	1	DHH1	XENLA	Q91610 xeno
62	7	2.5	398	1	DHH2	XENLA	Q91611 xeno
63	7	2.5	402	1	SELP	BOVIN	P49907 bos
64	7	2.5	412	1	PGKP	ALCEU	P50320 alca
65	7	2.5	413	1	PGKC	ALCEU	P50319 alca
66	7	2.5	416	1	NH59	CABEL	Q9txj1 caen
67	7	2.5	418	1	CP16	RAT	P09006 ratt
68	7	2.5	419	1	ENO	PYRAE	Q8zve7 pyro
69	7	2.5	419	1	ENK	RALSO	Q8ylw6 rals
70	7	2.5	423	1	YJ54	YEAST	P47130 sacc
71	7	2.5	424	1	YP3	MOUSE	P10761 mus
72	7	2.5	428	1	SYH	CHLMU	Q9pjj9 chia
73	7	2.5	455	1	PHR	STRGR	P12768 stre
74	7	2.5	461	1	PUCG	RHOCA	P23462 rhod
75	7	2.5	461	1	Y608	HAEIN	Q57486 haem
76	7	2.5	464	1	SOX8	MOUSE	Q04886 mus
77	7	2.5	467	1	D4DR	HUMAN	P21917 homo
78	7	2.5	483	1	GLME	CLOCC	P80077 clos
79	7	2.5	485	1	GLME	CLOTT	Q05509 clos
80	7	2.5	505	1	PDI	HUMIN	P55059 humi
81	7	2.5	512	1	LNT	ECOLI	P23910 esch
82	7	2.5	512	1	LNT	SALTY	O87576 balm
83	7	2.5	518	1	ASB3	HUMAN	Q9y575 homo
84	7	2.5	525	1	ASB3	MOUSE	Q9wv72 mus
85	7	2.5	547	1	RM56	HUMAN	P83111 homo
86	7	2.5	551	1	RM56	MOUSE	Q9ep89 mus
87	7	2.5	600	1	S133	MOUSE	Q9ly63 mus
88	7	2.5	600	1	S133	RAT	Q9z025 ratt
89	7	2.5	653	1	MUTL	VIBCH	Q9kv13 vibr
90	7	2.5	720	1	PPTA	PSEAE	P42512 vibr
91	7	2.5	758	1	CSTA	MYCTU	P95095 myco
92	7	2.5	760	1	PO21	XENLA	P16143 xeno
93	7	2.5	810	1	NELI	HUMAN	Q92832 homo
94	7	2.5	810	1	NELI	RAT	Q62919 ratt
95	7	2.5	854	1	KDPD	RATRA	O34971 ratt
96	7	2.5	853	1	B3A4	RAT	Q8k4v2 ratt
97	7	2.5	955	1	B3A4	RABIT	Q9gkv1 oryc
98	7	2.5	983	1	B3A4	HUMAN	Q96q31 oryc
99	7	2.5	1072	1	ITAC	CHICK	P26007 gall
100	7	2.5	1076	1	IF3A	CABEL	P34339 caen

ALIGNMENTS

RESULT 1

STANDARD; PRT; 249 AA.

Rel. 41, Created)

Rel. 41, Last sequence update)

Rel. 42, Last annotation update)

s factor ligand superfamily member 12 (TNF-related weak apoptosis) (TWEAK) (APO3 ligand).

03L OR DR3LG.

(Human)

Chordata; Craniata; Vertebrata; Euteleostomi; Hetera; Primates; Catarrhini; Hominidae; Homo.

06;

1 N.A., AND N-TERMINUS OF SOLUBLE FORM.

liver, and Tonsil;

015; PubMed=9405449;

ie Y., Bourdon P.R., Xu H., Hsu Y.-M., Scott H., Garcia I., Browning J.B.;

; secreted ligand in the tumor necrosis factor family that is apoptosis.";

1. 272:32401-32410(1997).

1 N.A.

Kidney;

0355; PubMed=9560343;

, Sheridan J.P., Pitti R.M., Brush J., Goddard A.,

on of a ligand for the death-domain-containing receptor

0:525-528(1998).

1 N.A.

0257; PubMed=12477932;

L., Feigold E.A., Grouse L.H., Derge J.G., Collins F.S., Wagner L., Shemen C.M., Schuler G.D., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K., Jordan H., Moore T., Max S.I., Wang J., Hsieh F., Marusina K., Farmer A., Rubin G.M., Hong L., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., J., Usdin T.B., Toshiyuki S., Carninci P., Prange C., Quellano N.A., Peters G.J., Abramson R.D., Mullahy S.J., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A., Ton E., Kettelman M., Madan A., Rodriguez S., Sanchez A., Adnan A., Young A.C., Shevchenko Y., Bouffard G.G., Touchman J.W., Green E.D., Dickson M.C., Grimwood J., Schmutz J., Myers R.M., S.N., Krzywinski M.I., Skalska U., Smailus D.E., Schein J.E., Jones S.J.M., Marra M.A.;

and initial analysis of more than 15,000 full-length cDNA sequences.";

Acad. Sci. U.S.A. 99:16899-16903(2002).

061; PubMed=10085077;

Hang Y.C., Lund J.K., Chen Y.-W., Leal J.A., Wiley S.R.;

is angiogenesis and proliferation of endothelial cells.";

1. 274:8455-8459(1999).

; Binds to FN14 and possibly also to TNFRSF12/APO3. Weak of apoptosis in some cell types. Mediates NF-kappaB on. May promote angiogenesis and the proliferation of al cells.

Homotrimer (Potential).

AR LOCATION: Type II membrane protein and secreted.

PECIFICITY: Highly expressed in adult heart, pancreas, muscle, brain, colon, small intestine, lung, ovary, spleen, lymph node, appendix and peripheral blood es. Low expression in kidney, testis, liver, placenta, id bone marrow. Also detected in fetal kidney, liver, brain.

CC -!- PTM: The soluble form derives from the membrane form

CC by proteolytic processing.

CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.

CC -!- CAUTION: Ref.3 sequence differs from that shown due to a frameshift in position 125.

CC -----

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CC -----

CC EMBL; AF030099; AAC51923.1; -

DR EMBL; AF055872; AAC39724.1; -

DR EMBL; BC019047; AAH19047.1; AUT_FRAME.

DR Genew; HGNC:11927; TNFSF12.

DR MIM; 602695; -

DR GO; GO:0005887; C:integral to plasma membrane; TAS.

DR GO; GO:0005102; F:receptor binding; TAS.

DR GO; GO:0006917; P:induction of apoptosis; TAS.

DR GO; GO:0007165; P:signal transduction; TAS.

DR InterPro; IPR006052; TNF family.

DR InterPro; IPR008983; TNF_like.

DR Pfam; PF00229; TNF; 1.

DR SMART; SM00207; TNF; 1.

DR PROSITE; PS00251; TNF_1; FALSE_NEG.

DR PROSITE; PS0049; TNF_2; 1.

DR Cytokine; Angiogenesis; Apoptosis; Transmembrane; Glycoprotein; signal-anchor.

KW CHAIN 1 249 TUMOR NECROSIS FACTOR LIGAND SUPERF

FT CHAIN 1 249 MEMBER 12, MEMBRANE FORM

FT CHAIN 94 249 TUMOR NECROSIS FACTOR LIGAND SUPERF

FT CHAIN 1 21 MEMBER 12, SECRETED FORM.

FT DOMAIN 22 42 CYTOPLASMIC (POTENTIAL).

FT TRANSMEM 22 42 SIGNAL-ANCHOR (TYPE-II MEMBRANE PRO

FT SITE 43 249 EXTRACELLULAR (POTENTIAL).

FT SITE 93 94 CLEAVAGE.

FT CARBOHYD 139 139 N-LINKED (GLCNAC...).

FT SEQUENCE 249 AA; 27216 MW; E660843361C28EBA CRC64;

SQ

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Best Local Similarity 100.0%; Pred. No. 2.9e-232;

Matches 249; Conservative 0; Mismatches 0; Indels 0;

QY 36 MAARRSQRRGRGEPGTALLVPLALGIGLALACIGLLAVVSLGSRASLSAQEP

Db 1 MAARRSQRRGRGEPGTALLVPLALGIGLALACIGLLAVVSLGSRASLSAQEP

QY 96 VAEEEDDPSELPQTEESQDPAPFLNLRPRRSPAPGKTRARRAIAAHVEVHP

Db 61 VAEEEDDPSELPQTEESQDPAPFLNLRPRRSPAPGKTRARRAIAAHVEVHP

QY 156 GAQAGVDGTSGWEERINSSPLRYNRQIGEFIVTRAGLYLYCQVHFDGKAV

Db 121 GAQAGVDGTSGWEERINSSPLRYNRQIGEFIVTRAGLYLYCQVHFDGKAV

QY 216 LLDVGVIALRCLEFSAATAASLGPQLRCQVSGLLALRPOSSLRIRTPWAHLK

Db 181 LLDVGVIALRCLEFSAATAASLGPQLRCQVSGLLALRPOSSLRIRTPWAHLK

QY 276 TYFGLFQVH 284

Db 241 TYFGLFQVH 249

RESULT 2

TN12_MOUSE

ID TN12_MOUSE STANDARD; PRT; 225 AA.

AC OS4907; Q9CTP2;

DT 28-FEB-2003 (Rel. 41, Created)

rel. 41, last sequence update)
 rel. 41, last annotation update)
 s factor ligand superfamily member 12 (TNF-related weak
 ptosis) (TWEAK) (fragment).
 (Mouse).
 azca; Chordata; Craniata; Vertebrata; Euteleostomi;
 eria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 090;
 N.A.
 real macrophage;
 il5; PubMed=9405449;
 a Y., Bourdon P.R., Xu H., Hsu Y.-M., Scott H.,
 arcia I., Browning J.L.;
 secreted ligand in the tumor necrosis factor family that
 s apoptosis.;
 : 272:32401-32410(1997).
 }-225 FROM N.A.
 J; TISSUE=Retina;
 60; PubMed=11217851;
 egawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
 ira A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
 wa M., Nishi K., Kiyosawa H., Kondo S., Yamana I.,
 aki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
 suda H.A., Ashburner M., Batalov S., Casavant T.,
 . Gaasterland T., Gissi C., King B., Kochiwa H.,
 is S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
 Staabli F., Suzuki R., Tomita M., Wagner L., Washio T.,
 io T., Furuno M., Anon H., Baldarelli R., Barsh G.,
 elli D., Bojunga N., Carninci P., de Bonaldo M.F.,
 f., Bult C., Fletcher C., Fujita M., Gariboldi M.,
 Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
 thionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 ng B., Ringwald M., Rodriguez I., Sakamoto N.,
 o K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
 o-oka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
 A., Yoshida K., Hasegawa Y., Kawaji H., Kobataki S.,
 ;
 notation of a full-length mouse cDNA collection.";
 }-690(2001).
 i binds to FN14 and possibly also to TNFRSF12/AP03. Weak
 i apoptosis in some cell types. Promotes angiogenesis and
 eration of endothelial cells. Mediates NF-kappaB
 l (By similarity).
 lonotrimer (Potential).
 AR LOCATION: Type II membrane protein and secreted (By
).
 IFFICITY: Widely expressed.
 oluble form is produced from the membrane form by
 C processing (By similarity).
 : Belongs to the tumor necrosis factor family.

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 profit institutions as long as its content is in no way
 his statement is not removed. Usage by and for commercial
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 ail to license@isb-sib.ch).

); AAC53517.1; -;
); BAB32249.1; -;
 259; Tnfsf12.
 06052; TNF family.
 08983; TNF_like.
); TNF; 1.
); TNF; 1.
 251; TNF_1; FALSE_NEG.
 049; TNF_2; 1.
 ogenesis; Apoptosis; Transmembrane; Glycoprotein;

FT NON TER 1 1 TUMOR NECROSIS FACTOR LIGAND SUPERFA
 FT CHAIN <1 225 MEMBER 12, MEMBRANE FORM.
 FT CHAIN 70 225 TUMOR NECROSIS FACTOR LIGAND SUPERFA
 FT CHAIN 70 225 MEMBER 12, SECRETED FORM (BY SIMILAR
 FT TRANSMEM <1 21 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROT
 FT DOMAIN 22 225 EXTRACELLULAR (POTENTIAL).
 FT SITE 69 70 CLEAVAGE (BY SIMILARITY).
 FT DISULFID 167 186 POTENTIAL.
 FT CARBOHYD 115 115 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 225 AA; 24781 MW; 90C412CC0480659B CRC64;
 Query Match 11.3%; Score 32; DB 1; Length 225;
 Best Local Similarity 100.0%; Pred. No. 4.3e-23;
 Matches 32; Conservative 0; Mismatches 0; Indels 0; G
 QY 139 RRAIAAHYEVHPRPGDGAQAGVDGTSGWEE 170
 D 80 RRAIAAHYEVHPRPGDGAQAGVDGTSGWEE 111
 RESULT 3
 ID YFBW ECOLI STANDARD; PRT; 111 AA.
 AC Q47377;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hypothetical protein yfbw.
 GN YFBW OR B2257.1.
 OS Escherichia coli.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
 OC Enterobacteriaceae; Escherichia.
 OC NCBI_TaxID=562;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=K12 / MG1655;
 RC MEDLINE=97426617; PubMed=9278503;
 RA Blattner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V
 RA Riley M., Collado-Vides J., Glaeser J.D., Rode C.K., Mayhew G.F.,
 RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
 RA Mau B., Shao Y.;
 RT "The complete genome sequence of Escherichia coli K-12.";
 RL Science 277:1453-1474(1997).
 RN [2]
 RP SEQUENCE OF 81-111 FROM N.A.
 RC STRAIN=K12;
 RC MEDLINE=96186953; PubMed=8626063;
 RA Sharma V., Hudepeth M.E., Megathathan R.;
 RT "Menquinone (vitamin K2) biosynthesis: localization and
 RT characterization of the menE gene from Escherichia coli.";
 RL Gene 168:43-48(1996).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a colla
 CC between the Swiss Institute of Bioinformatics and the EMBL out
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; AB000315; -; NOT ANNOTATED_CDS.
 DR EMBL; L35031; AAB04895.1; -;
 DR EcoGene; EGI4344; yfbw.
 DR InterPro; IPR000620; DUF6.
 DR Pfam; PF00892; DUF6; 1.
 KM Hypothetical protein; Transmembrane; Complete proteome.
 FT TRANSMEM 36 56 POTENTIAL.
 FT TRANSMEM 62 82 POTENTIAL.
 FT TRANSMEM 88 108 POTENTIAL.
 SQ SEQUENCE 111 AA; 12192 MW; 7CFA06D75DA33D69 CRC64;

3.5%; Score 10; DB 1; Length 111;
 Identity 100.0%; Pred. No. 0.038; 0; Indels 0; Gaps 0;
 Conservative 0; Mismatches 0; Indels 0; Gaps 0;

ALACGL 72
 |||||
 ALACGL 49

STANDARD; PRT; 733 AA.

(Rel. 33, Created)
 (Rel. 33, Last sequence update)
 (Rel. 42, Last annotation update)
 I P700 chlorophyll A apoprotein A2 (PsaB) (PSI-B).
 sensis (Marine centric diatom).

cramenophiles; Bacillariophyta; Coscinodiscophyceae;
 ycidae; Eupodiscaceae; Eupodiscaceae; Odontella.
 339;

M N.A.
 .. Stoebe B., Schaffran I., Kroth-Pancic P., Freier U.;
 last genome of a chlorophyll a+c-containing alga,
 sensis"; J. Phycol. 13:336-342(1995).

: PsaA and psaB bind P700, the primary electron donor of
 tem I (PSI), as well as the electron acceptors A0, A1 and
 is a plastocyanin/cytochrome c6-ferredoxin oxidoreductase,
 ng photonic excitation into a charge separation, which
 s an electron from the donor P700 chlorophyll pair to the
 copically characterized acceptors A0, A1, FX, FA and FB in
 ized P700 is reduced on the luminal side of the
 d membrane by plastocyanin or cytochrome c6.

: P700 is a chlorophyll A/chlorophyll A' dimer, A0 is one
 Chlorophyll A, A1 is one or both phylloquinones and FX is
 4Fe-4S iron-sulfur center (By similarity).
 The psaA/B heterodimer binds the P700 chlorophyll special
 subsequent electron acceptors. PSI consists of a core
 complex that captures photons, and an electron transfer
 at converts photonic excitation into a charge separation.
 ytic PSI reaction center is composed of at least 11
 (By similarity).

LAR LOCATION: Integral membrane protein. Chloroplast
 d membrane (By similarity).
 TY: Belongs to the psaA/psaB family.

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 mail to license@isb-sib.ch).

; CAA91749.1; -.

S78376.

; IJBO.

482; -; 1.

R006244; PsaB.

R001280; PSI_PsaA/B.

3; psaA_psaB; 1.

257; PHOTOSYSAAB.

GRO1336; PsaB; 1.

0419; PHOTOSYSTEM I_PSAAB; 1.

electron transport; Photosynthesis; Thylakoid;

I; Chlorophyll; Metal-binding; Iron; Magnesium;

4Fe-4S; Transmembrane; Chloroplast.

46

I (POTENTIAL).

FT TRANSMEM 134 157
 FT TRANSMEM 174 198
 FT TRANSMEM 272 230
 FT TRANSMEM 329 352
 FT TRANSMEM 368 394
 FT TRANSMEM 416 438
 FT TRANSMEM 516 534
 FT TRANSMEM 574 595
 FT TRANSMEM 642 664
 FT TRANSMEM 706 726
 FT METAL 558
 FT METAL 567 567
 FT METAL 653 653
 FT METAL 661 661
 FT BINDING 669 669
 FT BINDING 670 670
 SQ SEQUENCE 733 AA; 82103 MW; 13439AF1E441BBF7 CRC64;

Query Match 3.2%; Score 9; DB 1; Length 733;

Best Local Similarity 100.0%; Pred. No. 1.8;

Matches 9; Conservative 0; Mismatches 0; Indels 0;

QY 63 LGLALACLG 71

Db 333 LGLALACLG 341

RESULT 5

PSAB CVACA

ID PSAB CVACA STANDARD; PRT; 734 AA.

AC Q9TLQ6;

DT 16-OCT-2001 (Rel. 40, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Photosystem I P700 chlorophyll A apoprotein A2 (PsaB) (PSI-B).

GN PSAB

OS Cyanidium caldarium.

OG Chloroplast.

OC Eukaryota; Rhodophyta; Bangiophyceae; Porphyridiales; Porphyridi

OC Cyanidium.

OX NCBI_TaxID=2771;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=RK-1;

RX MEDLINE=20496959; PubMed=11040290;

RA Gloeckner G., Rosenthal A., Valentin K.-U.;

RT "The structure and gene repertoire of an ancient red algal plast

genome".

RL J. Mol. Evol. 51:382-390(2000).

CC -!- FUNCTION: PsaA and psaB bind P700, the primary electron dono

CC Photosystem I (PSI), as well as the electron acceptors A0, A

CC FX. PSI is a plastocyanin/cytochrome c6-ferredoxin oxidoredu

CC converting photonic excitation into a charge separation, whi

CC transfers an electron from the donor P700 chlorophyll pair t

CC spectroscopically characterized acceptors A0, A1, FX, FA and

CC turn. Oxidized P700 is reduced on the luminal side of the

CC thylakoid membrane by plastocyanin or cytochrome c6.

CC -!- COFACTOR: P700 is a chlorophyll A/chlorophyll A' dimer, A0 i

CC or more chlorophyll A, A1 is one or both phylloquinones and

CC a shared 4Fe-4S iron-sulfur center (By similarity).

CC -!- SUBUNIT: The psaA/B heterodimer binds the P700 chlorophyll s

CC pair and subsequent electron acceptors. PSI consists of a co

CC antenna complex that captures photons, and an electron trans

CC chain that converts photonic excitation into a charge separa

CC The eukaryotic PSI reaction center is composed of at least 1

CC subunits (By similarity).

CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Chloroplast

CC Thylakoid membrane (By similarity).

CC -!- SIMILARITY: Belongs to the psaA/psaB family.

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 or send an email to license@isb-sib.ch).

AAFL12881.1; -
 LUBO.
 12; -; 1
 06244; PsaB.
 01280; PSI_PsaA/B.
 psaA_psaB; 1.
 7; PHOTOSYSAAB.
 01336; PsaB; 1.
 19; PHOTOSYSTEM I PSAAB; 1.
 Chlorophyll; Photosynthesis; Thylakoid;
 Chlorophyll; Metal-binding; Iron; Magnesium;
 Fe-4S; Transmembrane; Chloroplast.
 6 69
 I (POTENTIAL).
 15 158
 II (POTENTIAL).
 75 199
 III (POTENTIAL).
 73 291
 IV (POTENTIAL).
 10 353
 V (POTENTIAL).
 19 395
 VI (POTENTIAL).
 7 439
 VII (POTENTIAL).
 7 535
 VIII (POTENTIAL).
 75 596
 IX (POTENTIAL).
 13 665
 X (POTENTIAL).
 7 727
 XI (POTENTIAL).
 19 559
 IRON-SULFUR (4FE-4S) (SHARED WITH DIMERIC
 PARTNER) (BY SIMILARITY).
 18 568
 IRON-SULFUR (4FE-4S) (SHARED WITH DIMERIC
 PARTNER) (BY SIMILARITY).
 14 654
 MAGNESIUM (CHLOROPHYLL-A B1 AXIAL LIGAND;
 P700 SPECIAL PAIR) (BY SIMILARITY).
 12 662
 MAGNESIUM (CHLOROPHYLL-A B3 AXIAL LIGAND)
 (BY SIMILARITY).
 10 670
 CHLOROPHYLL-A B3 (BY SIMILARITY).
 1 671
 PHYLOQUINONE B (BY SIMILARITY).
 1; AA; 82359 MW; 4496AA2AE59CA9B9 CRC64;

3.2%; Score 9; DB 1; Length 734;
 identity 100.0%; Pred. No. 1.8;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 AALG 71
 AALG 342

STANDARD; PRT; 179 AA.
 Rel. 35, Created)
 Rel. 35, Last sequence update)
 Rel. 41, Last annotation update)
 15 kDa subunit precursor (G3-ADH subunit III).
 oxydase (Gluconobacter suboxydase).
 eobacteria; Alphaproteobacteria; Rhodospirillales;
 ae; Gluconobacter.
 2;

N.A., AND SEQUENCE OF 26-40.
 328;
 225; PubMed=9055427;
 Inouchi S.;
 ion of the genes encoding the three-component membrane-
 dehydrogenase from Gluconobacter suboxydase and their

expression in Acetobacter pasteurianus.";
 Appl. Environ. Microbiol. 63:1131-1138 (1997).
 CC -!- FUNCTION: NOT ESSENTIAL FOR ALCOHOL DEHYDROGENASE ACTIVITY.
 CC -!- SUBUNIT: HETEROTRIMER (DEHYDROGENASE, CYTOCHROME AND PROTEIN
 CC ADHS). THAT FORMS THE ALCOHOL DEHYDROGENASE COMPLEX.
 CC -!- SUBCELLULAR LOCATION: Membrane-bound, facing the periplasmic
 CC (potential).
 CC

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 or send an email to license@isb-sib.ch).

EMBL; D86440; BAA19756.1; -
 Membrane; Periplasmic; Signal; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 24 POTENTIAL.
 FT CHAIN 25 179 ALCOHOL DEHYDROGENASE 15 kDa SUBUNIT
 FT MOD RES 25 25 PYRROLIDONE CARBOXYLIC ACID.
 SQ SEQUENCE 179 AA; 19943 MW; F6AF243656B3CC66 CRC64;

Query Match 2.8%; Score 8; DB 1; Length 179;
 Best Local Similarity 100.0%; Pred. No. 4.9;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G.

QY 59 LALGLGLA 66
 DB 11 LALGLGLA 18

RESULT 7
 Y304 BRUME
 ID Y304 BRUME STANDARD; PRT; 220 AA.
 AC CAYD73;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hypothetical protein BMEI10304.
 GN BMEI10304
 OS Brucella melitensis.
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
 OC Brucellaceae; Brucella.
 OX NCBI_TaxID=29459;
 RN 1;
 RP SEQUENCE FROM N.A.
 RC STRAIN=16M / ATCC 23456 / Biotype 1;
 RX MEDLINE=20020109; PubMed=11756688;
 RA DelVecchio V.G., Kaprat V., Redkar R.J., Patra G., Mijer C., Lo
 Ivanova N., Anderson I., Bhattacharya A., Lykidis A., Reznik G.,
 Jablonski L., Larsen N., D'Souza M., Bernal A., Mazur M., Goltzma
 Selkov E., Elzer P.H., Hagius S., O'Callaghan D., Letesson J.-J.,
 Haselkorn R., Kyriades N., Overbeek R.;
 RT The genome sequence of the facultative intracellular pathogen
 Brucella melitensis.";
 RT Brucella melitensis.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:443-448 (2002).
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
 CC -!- SIMILARITY: Belongs to the UPF0191 family.
 CC

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EMBL; AE009669; AAL53546.1; -
 DR PIR; AG3547; AG3547.
 DR HAMAP; MF 01207; -; 1.
 DR InterPro; IPR007916; UPF0191.
 DR Pfam; PF05252; UPF0191; 1.
 DR

06:25:21 2004

us-09-245-198a-4.oligo.rsp

protein; Transmembrane; Complete proteome.

20 39 POTENTIAL.
54 72 POTENTIAL.
85 104 POTENTIAL.
124 146 POTENTIAL.
153 175 POTENTIAL.
179 198 POTENTIAL.
20 AA; 24815 MW; 182C0244743B17FA CRC64;
2.8%; Score 8; DB 1; Length 220;
arity 100.0%; Pred. No. 5.8;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

/PLAL 61

/PLAL 139

STANDARD; PRT; 220 AA.

Rel. 43, Created)

Rel. 43, Last sequence update)

Rel. 43, Last annotation update)

protein BRA0991.

Neobacteria; Alphaproteobacteria; Rhizobiales;

Brucella.

1461;

1 N.A.

1 Biovar 1;

741; PubMed-12271122;

Seshadri R., Nelson K.E., Eisen J.A., Heidelberg J.F.,

Johnson R.J., Umayam L., Brinkac L.M., Beanan M.J.,

Deboy R.T., Durkin A.S., Kolonay J.F., Madupu R.,

Ayodeji B., Kraul M., Shetty J., Malek J., Van Aken S.E.,

Tetelin H., Gill S.R., White O., Salzberg S.L.,

Lindler L.E., Halling S.M., Boyle S.M., Fraser C.M.,

antis genome reveals fundamental similarities between

ant pathogens and symbionts."

Acad. Sci. U.S.A. 99:13148-13153(2002).

AR LOCATION: Integral membrane protein (Potential).

Y: Belongs to the UPF0191 family.

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12; AAN34160.1; -.

107; -; 1.

1007916; UPF0191.

1; UPF0191; 1.

protein; Transmembrane; Complete proteome.

20 39 Potential.

54 72 Potential.

85 104 Potential.

124 146 Potential.

153 175 Potential.

179 198 Potential.

20 AA; 24796 MW; AC2C060433169497 CRC64;

2.8%; Score 8; DB 1; Length 220;

arity 100.0%; Pred. No. 5.8;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 54 ALLVPLAL 61

DB 132 ALLVPLAL 139

RESULT 9

MSHR_PANTR

ID_MSHR_PANTR STANDARD; PRT; 317 AA.

AC Q9TUK4; O864L1;

DT 16-OCT-2001 (Rel. 40, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DT 15-MAR-2004 (Rel. 43, Last annotation update)

DE Melanocyte stimulating hormone receptor (MSH-R) (Melanotropin

DE receptor) (Melanocortin-1 receptor) (MCI-R).

GN MCI-R.

OS Pan troglodytes (Chimpanzee).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.

OX NCBI_TaxID=9598;

RN [1]

RP SEQUENCE FROM N.A.

RA Rees J.L., Harding R.M., Healy E., Jackson I.J., Ray A.J., Ellis

RA Flanagan N., Todd C., Dixon C., Matthews J.N., Sajantila A.,

RA Birch-Machin M.A.;

RT "Chimpanzee melanocortin 1 sequence."

RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=Isolate 3;

RX MEDLINE=22572539; PubMed=12687585;

RA Mundy N.I., Kelly J.;

RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, in

RT primates."

RL Am. J. Phys. Anthropol. 121:67-80(2003).

CC -!- FUNCTION: Receptor for MSH (alpha, beta and gamma) and ACTH.

CC activity of this receptor is mediated by G proteins which act

CC adenylylate cyclase (By similarity).

CC -!- SUBCELLULAR LOCATION: Integral membrane protein.

CC -!- SIMILARITY: Belongs to family 1 of G-protein coupled recepto.

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CC between the Swiss Institute of Bioinformatics and the EMBL out-

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CC use by non-profit institutions as long as its content is in

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5 315 S-palmitoyl cysteine (Potential).
AA; 34699 MW; 5615D2146E1D247F CRC64;
rity 2.8%; Score 8; DB 1; Length 317;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
IAA 144
IAA 167

STANDARD; PRT; 379 AA.

rel. 35, Created)
rel. 35, Last sequence update)
rel. 43, Last annotation update)
the dehydrogenase beta chain (EC 1.2.1.2).
jannaschii.
rchaeta; Methanococci; Methanococcales;
ccaceae; Methanocaldococcus.
0;
N.A.
DSM 2661 / ATCC 43067;
99; PubMed=6688087;
te O.; Olsen G.J.; Zhou L.; Fleischmann R.D.;
lake J.A.; FitzGerald L.M.; Clayton R.A.; Gocayne J.D.;
Dougherty B.A.; Tomb J.-F.; Adams M.D.; Reich C.I.;
Kirkness E.F.; Weinstock K.G.; Merrick J.M.; Glodek A.;
oghaen N.S.M.; Weidman J.F.; Fuhrmann J.L.; Nguyen D.;
Kelley J.M.; Peterson J.D.; Sadow P.W.; Hanna M.C.;
oberts K.M.; Hurst M.A.; Kaine B.P.; Borodovsky M.;
raser C.M.; Smith H.O.; Woese C.R.; Venter J.C.;
me sequence of the methanogenic archaeon, Methanococcus
58-1073(1996).
ACTIVITY: Formate + NAD(+) = CO(2) + NADH.
Binds 2 4Fe-4S clusters (Probable).
imer of alpha and beta chains (By similarity).
: The iron-sulfur centers are similar to those of
type 4Fe-4S ferredoxins.
: STRONG, TO THE BETA SUBUNIT OF M.THERMOAUTOTROPHICUM

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il to license@isb-sib.ch).

AAB37985.1; -
64300.

01450; 4Fe4S ferredoxin.
07525; FrhB_FdhB_C.
07516; FrhB_FdhB_N.
fer4; 1.
FrhB_FdhB_C; 1.
FrhB_FdhB_N; 1.
98; 4FE4S_FERREDOXIN; 2.
rotein; Oxidoreductase; NAD; Electron transport;
Fe-4S; Complete proteome.
0 280 IRON-SULFUR (4FE-4S) (BY SIMILARITY).
3 283 IRON-SULFUR (4FE-4S) (BY SIMILARITY).
6 286 IRON-SULFUR (4FE-4S) (BY SIMILARITY).
0 290 IRON-SULFUR (4FE-4S) (BY SIMILARITY).
0 330 IRON-SULFUR (4FE-4S) (BY SIMILARITY).

FT METAL 333 333 IRON-SULFUR (4FE-4S) (BY SIMILARITY)
FT METAL 336 336 IRON-SULFUR (4FE-4S) (BY SIMILARITY)
FT METAL 340 340 IRON-SULFUR (4FE-4S) (BY SIMILARITY)
SQ SEQUENCE 379 AA; 43014 MW; 9C257CCAD5547F5A CRC64;
Query Match 2.8%; Score 8; DB 1; Length 379;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 217 LVDGVLLAL 224
Db 35 LVDGVLLAL 42

RESULT 11
ZP3 MESAU STANDARD; PRT; 422 AA.
ID_ZP3 MESAU
AC P23491;
DT 01-NOV-1991 (Rel. 20, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 30-MAY-2000 (Rel. 39, Last annotation update)
DE Zona pellucida sperm-binding protein 3 precursor (Zona pellucida
glycoprotein ZP3) (Sperm receptor) (Zona pellucida protein C).
GN ZP3.
OS Mesocricetus auratus (Golden hamster).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae;
OC Mesocricetus.
OX NCBI_TaxID=10036;
RN [1]
SEQUENCE FROM N.A.
RP TISSUE=Ovary;
RC MEDLINE=91078540; PubMed=2257975;
RA Kinloch R.A.; Ruiz-Seller B.; Wassarman P.M.;
RT "Genomic organization and polypeptide primary structure of zona
pellucida glycoprotein HZP3, the hamster sperm receptor.";
RL Dev. Biol. 142:414-421(1990).
CC -!- FUNCTION: Functions as a sperm-receptor. It is responsible for
sperm-adhesion to the zona pellucida, and may contribute to ti
species-specificity of the insemination.
CC -!- SUBUNIT: ZP3 FORMS WITH ZP1 AND ZP2 THE ZONA PELLUCIDA, IN
WHICH ZP2 AND ZP3 COMPLEX INTO COPOLYMERS CROSS-LINKED BY ZP1
CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Extracellular
matrix.
CC -!- TISSUE SPECIFICITY: Oocytes.
CC -!- DEVELOPMENTAL STAGE: GROWING OOCYTES.
CC -!- PTM: Sulfated glycoprotein with O-linked oligosaccharides.
CC -!- SIMILARITY: Contains 1 ZP domain.

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EMBL; M63629; AAA37079.1;
InterPro; IPR001507; Endoglin/CD105.
Fram; PF00100; zona_pellucida; 1.
PRINTS; PRO0023; ZPELLUCIDA.
SMART; SM00241; ZP; 1.
DR PROSITE; PS00682; ZP_DOMAIN; 1.
KW Glycoprotein; Signal; Sulfation; Sperm; Receptor; Transmembrane;
Extracellular matrix.
FT SIGNAL 1 22
FT CHAIN 23 422
FT DOMAIN 23 386
FT TRANSMEM 387 407
FT DOMAIN 408 422
FT DOMAIN 45 306
FT DOMAIN 119 158
FT DOMAIN 208 257
POTENTIAL.
ZONA PELLUCIDA SPERM-BINDING PROTEIN
EXTRACELLULAR (POTENTIAL).
POTENTIAL.
CYTOPLASMIC (POTENTIAL).
ZP.
PRO-RICH.
PRO-RICH.

146 146 N-LINKED (GLCNAC. . .) (POTENTIAL).
 271 271 N-LINKED (GLCNAC. . .) (POTENTIAL).
 302 302 N-LINKED (GLCNAC. . .) (POTENTIAL).
 22 AA; 45827 MW, D0F95B7FFBE7E01 CRC64;

2.8%; Score 8; DB 1; Length 422;
 larity 100.0%; Pred. No. 10;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;

ILGIA 66

|||||

ILGIA 393

STANDARD; PRT; 576 AA.

(Rel. 32, Created)

(Rel. 32, Last sequence update)

(Rel. 41, Last annotation update)

p-binding protein cydC.

36.

teobacteria; Gammaproteobacteria; Pasteurellales;

ae; Haemophilus.

27;

4 N.A.

KW20 / ATCC 51907;

3630; PubMed:7542800;

3.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,

3., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,

Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,

Shirley R., Liu L.-I., Glodek A., Kelley J.M.,

, Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,

3., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,

itchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,

McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,

a random sequencing and assembly of Haemophilus influenzae

196-512(1995).

: SOMEHOW INVOLVED IN THE CYTOCHROME D BRANCH OF AEROBIC

ION. SEEMS TO BE A COMPONENT OF A TRANSPORT SYSTEM

LARITY).

AR LOCATION: Integral membrane protein. Inner membrane

3). Belongs to the ABC transporter family. MsbA subfamily.

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; AAC22811.1; --

; E64186.

; --

R003593; AAA ATPase.

R001140; ABC_TM transp.

R003439; ABC transporter.

4; ABC membrane; 1.

5; ABC_tran; 1.

3006; ABC transporter; 1.

32; AAA; 1.

3929; ABC TMIF; 1.

3211; ABC_TRANSPORTER_1; 1.

3893; ABC_TRANSPORTER_2; 1.

Transp.; transmembrane; Inner membrane;

ecome.

FT TRANSMEM 16 36 POTENTIAL.
 FT TRANSMEM 38 58 POTENTIAL.
 FT TRANSMEM 133 153 POTENTIAL.
 FT TRANSMEM 155 175 POTENTIAL.
 FT TRANSMEM 244 264 POTENTIAL.
 FT TRANSMEM 281 301 POTENTIAL.
 FT DOMAIN 338 374 ABC TRANSPORTER.
 FT NP_BIND 372 379 ATP (POTENTIAL).
 SQ SEQUENCE 576 AA; 64831 MW; A9ACD8B9B294B1B3 CRC64;

Query Match 2.8%; Score 8; DB 1; Length 576;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 58 PLALGLGL 65

Db 159 PLALGLGL 166

RESULT 13

GGT5 HUMAN

ID GGT5 HUMAN STANDARD; PRT; 586 AA.

AC P36269; Q96FC1; Q9UFM5;

DT 01-JUN-1994 (Rel. 29, Created)

DT 01-JUN-1994 (Rel. 29, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Gamma-glutamyltransferase 5 precursor (EC 2.3.2.2) (Gamma-glutamyltransferase 5) (Gamma-glutamyltransferase-like activi-

DE (GGT-rel).

GN GGT1A1 OR GGT5.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi

OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A. (ISOFORM 1).

RC TISSUE=Placenta;

RX MEDLINE=91296809; PubMed=1676842;

RA Heisterkamp N., Rajpert-De Meyts E., Uribe L., Forman H.J.,

RA Groffen J.,

RT "Identification of a human gamma-glutamyl cleaving enzyme relate but distinct from, gamma-glutamyl transpeptidase."

RL Proc. Natl. Acad. Sci. U.S.A. 88:6303-6307(1991).

RN [2]

RP SEQUENCE FROM N.A. (ISOFORM 2).

RC TISSUE=Petal kidney;

RA Blum H., Bauersachs S., Mewes H.-W., Gassenhuber J., Wiemann S.;

RL Submitted (SEP-1999) to the EMBL/GenBank/DBJ databases.

RN [3]

RP SEQUENCE FROM N.A.

RX MEDLINE=20057165; PubMed=10591208;

RA Dunham I., Hunt A.R., Collins J.E., Bruskewich R., Beare D.M.,

RA Clamp M., Smink L.J., Ainscough R., Almeida J.P., Babbage A.K.,

RA Bagguley C., Bailey J., Barlow K.F., Bates K.N., Beasley O.P.,

RA Bird C.P., Blakey S.B., Bridgman A.M., Buck D., Burgess J.,

RA Burrill W.D., Burton J., Carder C., Carter N.P., Chen Y., Clark I

RA Clegg S.M., Cobley V.E., Cole C.G., Collier R.E., Connor R.,

RA Conroy D., Corby N.R., Coville G.J., Cox A.V., Davis J., Dawson I

RA Dhani P.D., Dockree C., Dodsworth S.J., Durbin R.M., Ellington A

RA Evans K.L., Fey J.M., Fleming K., French L., Garner A.A.,

RA Gilbert J.G.R., Goward M.E., Grafham D.V., Griffiths M.N.D., Hall

RA Hall R.E., Hall-Tamlyn G., Heathcote R.W., Ho S., Holmes S.,

RA Hunt S.E., Jones M.C., Kerhaw J., Kimberley A.M., King A.,

RA Laird G.K., Langford C.F., Leversha M.A., Lloyd C., Lloyd D.M.,

RA Martyn I.D., Mashreghi-Mohammadi M., Matthews L.H., Mccann O.T.,

RA Mcclay J., McLaren S., McMurray A.A., Milne S.A., Mortimore B.J.

RA Odeil C.N., Pavitt R., Pearce A.V., Pearson D., Phillimore B.J.C

RA Phillips S.H., Plumb R.W., Ramsay H., Ramsey Y., Rogers L., Ross

RA Scott C.E., Sehra H.K., Skuce C.D., Smalley S., Smith M.L.,

RA Soderlund C., Spragon L., Steward C.A., Sulston J.E., Swann R.M.

RA Vaudin M., Wall M., Wallis J.M., Whiteley M.N., Willey D.B.,

RA Williams L., Williams S.A., Williamson H., Wilmer T.E., Wilming I

RA Wright C.L., Hubbard T., Bentley D.R., Beck S., Rogers J., Shimi:

Kawasaki K., Sasaki T., Asakawa S., Kudoh J., Shibuya K., Yoshiaki Y., Aoki N., Mitsuyama S., Chu L., Crabtree J., Deschamps S., Do A., Do T., Fu Y., Hu P., Hua A., Kenton S., Lai H., Lao H.I., Lin S.-P., Loh P., Malaj E., Nguyen T., Pan H., Qian Y., Ray L., Ren Q., Shaull S., Sloan D., Song L., Wang Z., White J., Willingham D., Wu H., Yao Z., Y., Chissos S., Murray J., Miller N., Mind P., nson D., Bemis G., Bentley D., Bradshaw H., Bourne S., Fulton L., Goela D., Graves T., Hawkins J., K., Latreille P., Layman D., Ozersky P., Rohlfing T., lker C., Wamley A., Wohlmann P., Pepin K., Nelson J., J.A., Hillier L.W., Mardis E., Waterston R., mel B.S., Shaikh T., Kurahashi H., Saitta S., mberland H.E., Johnson A., Wong A.C.C., Morrow B.E., Kim U.J., Shizuya H., Simon M.I., Dumanski J.P., edra D., Seroussi E., Francon I., Tapia I., Bruder C.E., Wilkerson P., Bodenteich A., Hartman K., Hu X., e L., Tiliahun Y., Wright H.;
ence of human chromosome 22."
3-495(1999).
N.A. (ISOFORM 1).
257; PubMed=12477932;
J., Feingold E.A., Grouse L.H., Derge J.G., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K., Jordan H., Moore T., Max S.I., Wang J., Heieh F., Marusina K., Farmer A.A., Rubin G.M., Hong L., Soares M.B., Bonaldo M.F., Casavant T.B., Scheetz T.E., Udén T.B., Toshiyuki S., Carninci P., Prange C., Ruellano N.A., Peters G.J., Abramson R.D., Mullahy S.J., Ewan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A., Con E., Kettman M., Madan A., Rodrigues S., Sanchez A., adan A., Young A.C., Shevchenko Y., Bouffard G.G., Touchman J.W., Green E.D., Dickson M.C., Grimwood J., Schmutz J., Myers R.M., S.N., Krzywinski M.I., Skalska U., Smallos D.E., Schein J.B., Jones S.J.M., Marra M.A.;
ad initial analysis of more than 15,000 full-length cdna sequences."
ad. Sci. U.S.A. 99:16899-16903(2002).
cleaves the gamma-glutamyl peptide bond of glutathione
s, but maybe not glutathione itself. Converts
le C4 (LTD4) to leukotriene D4 (LTD4).
ACTIVITY: (S-L-glutamyl)-peptide + an amino acid =
S-L-glutamyl-amino acid.
glutathione metabolism.
leukotriene metabolism; second step.
eterodimer composed of the light and heavy chains.
e site is located in the light chain (By similarity).
AR LOCARION: Type II membrane protein (By similarity).
VE PRODUCTS:
ernative splicing; Named isoforms=2;
36269-1; Sequence=Displayed;
36269-2; Sequence=VSP_008146;
experimental confirmation available;
y: Belongs to the gamma-glutamyltransferase family.
Ref.2 sequence differs from that shown due to a
t in position 446.
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CC EMBL; M64099; AAA58503.1; --
DR EMBL; AL117414; CAB55910.1; ALT FRAME.
DR EMBL; AP000354; -- NOT ANNOTATED_CDS.
DR EMBL; BC011362; AAH11362.1; --
DR PIR; A41125; A41125.
DR PIR; T17220; T17220.
DR MEROPS; T03.002; --
DR Genew; HGNC:4260; GGTAL1.
DR MIM; 137168; --
DR GO; GO:0016021; C: integral to membrane; TAS.
DR GO; GO:0003840; F: gamma-glutamyltransferase activity; TAS.
DR GO; GO:0006520; P: amino acid metabolism; TAS.
DR GO; GO:0006749; P: glutathione metabolism; TAS.
DR InterPro; IPR000101; Peptidase T3.
DR Pfam; PF01019; G: glutathione; 1.
DR PRINTS; PS0210; G: TRANSFERASE.
DR PROSITE; PS00462; G: GLUTAMYLTRANSFERASE; 1.
KW Glutathione biosynthesis; Leukotriene biosynthesis; Transferase;
KW Acyltransferase; Signal-anchor; Transmembrane; Zymogen; Glycoprot
KW Alternative splicing.
FT CHAIN 1 387
FT CHAIN 388 586
FT CHAIN (BY SIMILARITY).
FT CHAIN (BY SIMILARITY).
FT DOMAIN 1 8
FT TRANSMEM 9 29
FT SIGNAL-ANCHOR (TYPE-II MEMBRANE PROT
FT (POTENTIAL).
FT DOMAIN 30 586
FT CARBOHYD 98 98
FT CARBOHYD 204 204
FT CARBOHYD 303 303
FT CARBOHYD 347 347
FT CARBOHYD 535 535
FT CARBOHYD 550 550
FT VARSPLIC 101 132
FT Missing (in isoform 2).
FT CONFLICT 330 330
FT CONFLICT 408 408
FT CONFLICT 437 437
FT CONFLICT 586 AA; 62319 MW; 1BE543CB0934B16B CRC64;
SQ SEQUENCE 2.8%; Score 8; DB 1; Length 586;
Query Match Best Local Similarity 100.0%; Pred. No. 14;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G
QY 61 LGGLALA 68
DB 14 LGGLALA 21
RESULT 14
HSP3 OCTVU STANDARD; PRT; 24 AA.
AC P83215;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Sperm protamine P3 (Po3) (Fragment).
OS Octopus vulgaris (Octopus).
OC Eukaryota; Metazoa; Mollusca; Cephalopoda; Coleoidea; Neocoleoide
OC Octopodiformes; Octopoda; Incirrata; Octopodidae; Octopus.
OX NCBI_TaxID=6645;
RN [1]
RP SEQUENCE, FUNCTION, AND MASS SPECTROMETRY.
RC TISSUE=Sperm;
RA Gimenez-Bonafe P., Ribes E., Buesa C., Sautiere P., Kouach M.,
RA Ausio J., Kasinsky H.E., Chiva M.;
RT "Chromatin remodelling and protamines during spermiogenesis of Oc
RT vulgaris (Cephalopoda)."
RL J. Exp. Zool. 0:0-0(2001).
CC -!- FUNCTION: Protamines substitute for histones in the chromatin
sperm during the haploid phase of spermatogenesis. They compa

A into a highly condensed, stable and inactive

LAR LOCATION: Nuclear.

PECIFICITY: Testis.

CTROMETRY: MW=4389; METHOD=Electrospray.

36; C:nucleosome; NAS.

34; C:nucleus; NAS.

77; F:DNA binding; NAS.

01; P:chromosome organization and biogenesis (sen. . .; NAS.

76; P:mitotic chromosome condensation; NAS.

34; P:nucleosome assembly; NAS.

33; P:spermatogenesis; NAS.

protein; Nucleosome core; Spermatogenesis;

DNA condensation; Nuclear protein.

1 16 POLY-ARG.

24 24

4 AA; 3381 MW; 308E90ED9D2C9C9C CRC64;

2.5%; Score 7; DB 1; Length 24;

larity 100.0%; Pred.No. 7.7;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

3RRG 49

||||

3RRG 17

STANDARD; PRT; 30 AA.

(Rel. 41, Created)

(Rel. 41, Last sequence update)

(Rel. 41, Last annotation update)

ine P5 (Po5).

aris (Octopus).

stazoa; Mollusca; Cephalopoda; Coleoidea; Neocoleoidea;

es; Octopoda; Incirrata; Octopodidae; Octopus.

645;

NCTION, AND MASS SPECTROMETRY.

ife P., Ribes E., Buesa C., Sautiere P., Kouach M.,

sinsky H.E., Chiva M.;

emodelling and protamines during spermiogenesis of Octopus

phalopoda).";

0.0-0.0(2001).

: Protamines substitute for histones in the chromatin of

ring the haploid phase of spermatogenesis. They compact

A into a highly condensed, stable and inactive

LAR LOCATION: Nuclear.

PECIFICITY: Testis.

CTROMETRY: MW=3941; METHOD=Electrospray.

36; C:nucleosome; NAS.

34; C:nucleus; NAS.

77; F:DNA binding; NAS.

01; P:chromosome organization and biogenesis (sen. . .; NAS.

76; P:mitotic chromosome condensation; NAS.

34; P:nucleosome assembly; NAS.

33; P:spermatogenesis; NAS.

protein; Nucleosome core; Spermatogenesis;

DNA condensation; Nuclear protein.

2 15 POLY-ARG.

17 26

0 AA; 3943 MW; 14F1BC7E4D277049 CRC64;

2.5%; Score 7; DB 1; Length 30;

larity 100.0%; Pred.No. 9.3;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

3RRG 49

||||

Db

5 RRRGRRG 11

RESULT 16

HSPI_SAGIM

ID HSPI_SAGIM STANDARD; PRT; 49 AA.

DT 01-MAR-1992 (Rel. 21, Created)

DT 01-MAR-1992 (Rel. 21, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Sperm protamine P1 (Cysteine-rich protamine).

GN PRM1.

OS Saguinus imperator (Emperor tamarin).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi

OC Mammalia; Eutheria; Primates; Platyrrhini; Callitrichidae; Sagu

OX NCBI_TaxID=9491;

RN [1]

SEQUENCE FROM N.A.

RC TISSUE=Liver;

RX MEDLINE=92051332; PubMed=1840669;

RA Queralt R., Oliva R.;

RT "Protamine 1 gene sequence from the primate Saguinus imperator

isolated with PCR using consensus oligonucleotides.";

RL Nucleic Acids Res. 19:5786-5786(1991).

CC -!- FUNCTION: Protamines substitute for histones in the chromati

sperm DNA into a highly condensed, stable and inactive compl

-!- SUBUNIT: Cross-linked by interchain disulfide bonds around t

DNA-helix (By similarity).

-!- SUBCELLULAR LOCATION: Nuclear.

-!- TISSUE SPECIFICITY: Testis.

-!- SIMILARITY: Belongs to the protamine P1 family.

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DR EMBL; X61678; CAA43853.1; --

PIR; S22582; S22582.

DR InterPro; IPR000221; Protamine.P1.

DR Pfam; PF00260; Protamine.P1; 1.

DR PROSITE; PS00048; PROTAMINE.P1; 1.

KW Chromosomal protein; Nucleosome core; Spermatogenesis; DNA-bindi

Testis; DNA condensation; Nuclear protein.

FT INIT_MET 0 0

SQ SEQUENCE 49 AA; 6545 MW; 8389C403F5B207F6 CRC64;

Query Match 2.5%; Score 7; DB 1; Length 49;

Best Local Similarity 100.0%; Pred.No. 14;

Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 42 QRRRGR 48

|||||

Db 17 QRRRGR 23

RESULT 17

HSPI_DIDMA

ID HSPI_DIDMA STANDARD; PRT; 57 AA.

AC P35305;

DT 01-FEB-1994 (Rel. 28, Created)

DT 01-FEB-1994 (Rel. 28, Last sequence update)

DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE Sperm protamine P1.

GN PRM1.

OS Didelphis marsupialis virginiana (North American opossum), and

OC Monodelphis domestica (Short-tailed grey opossum).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi

OC Mammalia; Metatheria; Didelphimorphia; Didelphidae; Didelphis.

57, 13616;
 N.A.
 supialis;
 300; PubMed=8344286;
 Nishikawa S., Connor W., Dixon G.H.;
 tion of a marsupial sperm protamine gene and its
 com the North American opossum (*Didelphis*
 ';
 m. 215:63-72(1993).
 N.A.
 istics;
 51; PubMed=7700877;
 Gajewski C., Western M., Winkfein R.J., Dixon G.H.;
 logeny and evolution of marsupial protamine P1 genes.;
 Lond., B. Biol. Sci. 259:7-14(1995).
 Protamines substitute for histones in the chromatin of
 ing the haploid phase of spermatogenesis. They compact
 into a highly condensed, stable and inactive complex.
 AR LOCATION: Nuclear.
 SPECIFICITY: Testis.
 (: Belongs to the protamine P1 family.
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 ail to license@isb-sib.ch).

 AAA02812.1; -;
 CAA52193.1; -;
 AAA74612.1; -;
 334045.
 000221; Protamine P1.
 protamine P1; 1.
 148; PROTAMINE P1; 1.
 corein; Nucleosome core; Spermatogenesis; DNA-binding;
 ndensation; Nuclear protein.
 0
 0 BY SIMILARITY.
 AA; 7810 MW; 283715820214E52 CRC64;
 2.5%; Score 7; DB 1; Length 57;
 100.0%; Pred No. 16;
 0; Mismatches 0; Indels 0; Gaps 0;
 RG 49
 ||
 RG 40
 STANDARD; PRT; 115 AA.
 Rel. 40, Created)
 Rel. 40, Last sequence update)
 Rel. 42, Last annotation update)
 id protein Acp62F precursor.
 162.
 lanogaster (Fruit fly).
 :azoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 pterygota; Diptera; Brachycera; Muscomorpha;
 rosophilidae; Drosophila.
 ?;
 N.A., FUNCTION, AND TISSUE SPECIFICITY.
 -S; TISSUE=Male accessory gland;
 20; PubMed=9474779;
 Harada H.A., Bertram M.J., Stelick T.J., Kraus K.W.,
 ig Y.O., Neubaum D.M., Park M., Tram U.K.;

RT
 RT melanogaster.";
 RL Insect Biochem. Mol. Biol. 27:825-834(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkeley;
 RX MEDLINE=20196006; PubMed=107311132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.B., Richards S., Ashburner M., Henderson S.N
 RA Sutton G.G., Wortman J.R., Fandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L
 RA Abail J.F., Agbayani A., An H.-J., Andrews-Pfankoch C., Baldwin
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Bocham M.R., Bouck J., Brokstein P., Brottier P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dun
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischman
 RA Foster K., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodok A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D.A., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lasko P., Lei Y., Levitsky A.A., Li J.H., Li Z., Liang Y., Lin X.
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spadling A.C., Stapleton M., Strong R., Sun E.,
 RA Svirskaas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wassarman D.A., Weinstock G.M., Weissenbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of *Drosophila melanogaster*.";
 RL Science 287:2185-2195(2000).
 RN [3]
 RP SEQUENCE OF 7-111 FROM N.A.
 RC STRAIN=ZIM62H-12C, ZIM62H-16C, ZIM62H-28C, ZIM62H-30C, ZIM62H-34C
 RC ZIM62I-5C, ZIM62I-10C, ZIM62I-17C, ZIM62I-18C, and ZIM62I-53C;
 RX MEDLINE=20556153; PubMed=11102381;
 RA Begun D.J., Whitley P., Todd B.L., Waldrup-Dail H.M., Clark A.G.;
 RT "Molecular population genetics of male accessory gland proteins i
 RL *Drosophila*.";
 RL Genetics 156:1879-1888(2000).
 CC -I- FUNCTION: RESPONSIBLE FOR PHYSIOLOGICAL AND BEHAVIORAL CHANGE;
 CC MATED FEMALE FLIES. MAY CONTRIBUTE TO THE TOXICITY OF SEMINAL
 CC FLUID AND THE DECREASED LIFE-SPAN OF MATED FEMALES. MAY ALSO
 CC AFFECT NEUROMUSCULAR EVENTS AFTER MATING CONCERNING SPERM STO
 CC AND EGG RELEASE.
 CC -I- SUBCELLULAR LOCATION: Secreted (Probable).
 CC -I- TISSUE SPECIFICITY: Seminal fluid.
 CC -I- SIMILARITY: SOME, TO P.NIGRIVENTER TX2-6.
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 DR EMBL; U85763; AAB96387.1; -;
 DR EMBL; AB003475; AAF47683.1; -;

8; AAG35367.1; -
 9; AAG35368.1; -
 0; AAG35369.1; -
 1; AAG35370.1; -
 2; AAG35371.1; -
 3; AAG35372.1; -
 4; AAG35373.1; -
 5; AAG35374.1; -
 6; AAG35375.1; -
 7; AAG35376.1; -
 10020509; ACP62F.
 17; F:serine protease inhibitor activity; IDA.
 0; P:determination of adult life span; NAS.
 3002919; TIL_Cyrich.
 1; TIL; 1.
 anal.
 1 24 POTENTIAL.
 25 115 ACCESSORY GLAND PROTEIN ACP62F.
 34 88 TIL.
 5 AA; 12570 MW; 4326AA6F6C32291D CRC64;
 2.5%; Score 7; DB 1; Length 115;
 arity 100.0%; Pred.No.30;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 ELL 74
 ||||
 ELL 17
 STANDARD; PRT; 118 AA.
 Rel. 01, Created)
 Rel. 01, Last sequence update)
 Rel. 41, Last annotation update)
 ment ISS very hypothetical 12 kDa protein.
 coli.
 reobacteria; Gammaproteobacteria; Enterobacteriales;
 aceae; Escherichia.
 1 N.A.
 1653; PubMed=6269959;
 Gahn M.;
 de sequence of ISS from Escherichia coli.";
 .74(1981).
 1 N.A.
 1652; PubMed=6269958;
 van Bree M.P.;
 de sequence and protein-coding capability of the
 element ISS.";
 .63(1981).
 1 N.A.
 Duncan M., Allen E., Araujo R., Aparicio A., Chung E.,
 Ierspiel N., Hyman R., Kalman S., Komp C., Kurdi O.,
 Lew H., Lin D., Namath A., Oefner P., Roberts D.,
 (P-1996) to the EMBL/GenBank/DBJ databases.
 1 N.A.
 Mori H., Murayama N., Kataoka K., Yano M., Itoh T.,
 Inokuchi H., Maki T., Hatada E., Fukuda R.,
 Mizuno T., Makino K., Nakata A., Yura T., Sampei G.,
 (B-1996) to the EMBL/GenBank/DBJ databases.
 1 N.A.

RX MEDLINE=97251357; PubMed=9097039;
 RA Aiba H., Baba T., Fujita K., Hayashi K., Inada T., Isono K.,
 RA Itoh T., Kasai H., Kashimoto K., Kimura S., Kitakawa M.,
 RA Kitagawa M., Makino K., Miki T., Mizobuchi K., Mori H., Mori T.,
 RA Motomura K., Nakade S., Nakamura Y., Nishimoto H., Nishio Y.,
 RA Oshima T., Saito N., Sampei G., Seki Y., Sivasubram S.,
 RA Tagami H., Takeda J., Takemoto K., Takeuchi Y., Wada C.,
 RA Yamamoto Y., Horiuchi T.;
 RT "A 570-kb DNA sequence of the Escherichia coli K-12 genome
 RT corresponding to the 28.0-40.1 min region on the linkage map.";
 RL DNA Res. 3:363-377(1996).
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 CC or send an email to license@isb-sib.ch).
 CC EMBL; J01734; -; NOT ANNOTATED_CDS.
 DR EMBL; U70214; AAB08680.1; -
 DR EMBL; D83336; -; NOT ANNOTATED_CDS.
 DR EMBL; D90771; BAA14925.1; -
 DR EMBL; D90772; BAA14935.1; -
 DR EMBL; D90831; BAA15715.1; -
 DR EMBL; D90841; BAA15872.1; -
 DR EMBL; D90847; BAA15958.1; -
 DR EMBL; D90848; BAA15963.1; -
 DR PIR; B91483; IEEC5B.
 KW Hypothetical protein; Transposable element.
 SQ SEQUENCE 118 AA; 12270 MW; 348014FAC765058E CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 118;
 Best Local Similarity 100.0%; Pred.No.31;
 Matches 7; Conservative 0; Mismatches 0; Indels 0;
 QY 23 DGGAVRQ 29
 |||||
 DB 107 DGGAVRQ 113
 RESULT 20
 IL13 MOUSE
 ID IL13 MOUSE STANDARD; PRT; 131 AA.
 AC P20109.
 DT 01-FEB-1991 (Rel. 17, Created)
 DT 01-FEB-1991 (Rel. 17, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE Interleukin-13 precursor (IL-13) (T-cell activation protein P600
 GN IL13 OR IL-13.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mu-
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=89093958; PubMed=2521353;
 RA Brown K.D., Zurawski S.M., Mosmann T.R., Zurawski G.;
 RT "A family of small inducible proteins secreted by leukocytes are
 RT members of a new superfamily that includes leukocyte and
 RT fibroblast-derived inflammatory agents, growth factors, and
 RT indicators of various activation processes.";
 RL J. Immunol. 142:679-687(1989).
 CC -!- FUNCTION: CYTOKINE. INHIBITS INFLAMMATORY CYTOKINE PRODUCTION
 CC SYNERGIZES WITH IL2 IN REGULATING INTERFERON-GAMMA SYNTHESIS
 CC MAY BE CRITICAL IN REGULATING INFLAMMATORY AND IMMUNE RESPON-
 CC (BY SIMILARITY).
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- SIMILARITY: Belongs to the IL-4 / IL-13 family.
 CC This SWISS-PROT entry is copyright. It is produced through a col-
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 AAA40149.1; -.
 31TR.
 ; IL13.
 03634; Interleukin_13.
 01325; Interleukin_4_13.
 Interleukin_13; 1.
 ; IL4_13; 1.
 ; IL4_13; 1.
 138; INTERLEUKIN_4_13; 1.
 oprotein; Signal.
 1 21 BY SIMILARITY.
 2 131 INTERLEUKIN-13.
 3 79 BY SIMILARITY.
 4 93 BY SIMILARITY.
 5 42 N-LINKED (GLCNAC. . .) (POTENTIAL).
 6 52 N-LINKED (GLCNAC. . .) (POTENTIAL).
 7 75 N-LINKED (GLCNAC. . .) (POTENTIAL).
 AA; 14107 MW; 954F93F105713FED CRC64;

2.5%; Score 7; DB 1; Length 131;
 100.0%; Pred. No. 34;
 0; Mismatches 0; Indels 0; Gaps 0;

IG 71
 ||
 IG 15

STANDARD; PRT; 131 AA.

rel. 32, Created)
 rel. 32, Last sequence update)
 rel. 36, Last annotation update)
 precursor (IL-13) (T-cell activation protein P600).

cus (Rat).
 azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 eria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 16;

N.A.
 ; Dawley; TISSUE=Kidney cortex;
 38; PubMed=7916615;

it interleukin-13 (IL-13) cDNA and analysis of IL-13
 in experimental glomerulonephritis";
 Mys. Res. Commun. 197:612-618(1993).

CYTOKINE INHIBITS INFLAMMATORY CYTOKINE PRODUCTION.
 WITH IL2 IN REGULATING INTERFERON-GAMMA SYNTHESIS
 TICAL IN REGULATING INFLAMMATORY AND IMMUNE RESPONSES
 ARITY).

AR LOCATION: Secreted.

); Belongs to the IL-4 / IL-13 family.

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 AAA16478.1; -.
 152290.

DR HSP; P35225; 31TR.
 DR InterPro; IPR003634; Interleukin_13.
 DR InterPro; IPR001325; Interleukin_4_13.
 DR Pfam; PF03487; Interleukin_13; 1_
 DR ProDom; PD015987; Interleukin_13; 1.
 DR SMART; SM00190; IL4_13; 1.
 DR PROSITE; PS00838; INTERLEUKIN_4_13; 1.
 KW Cytokine; Glycoprotein; Signal.
 FT SIGNAL 1 21 BY SIMILARITY.
 FT CHAIN 22 131 INTERLEUKIN-13.
 FT DISULFID 52 80 BY SIMILARITY.
 FT DISULFID 68 94 BY SIMILARITY.
 FT CARBOHYD 42 42 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 53 53 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 76 76 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 121 121 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 131 AA; 14093 MW; E5008CAB8DE8C201 CRC64;

Query Match 2.5%; Score 7; DB 1; Length 131;
 Best Local Similarity 100.0%; Pred. No. 34;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 65 LALACLG 71
 |||||
 DB 9 LALACLG 15

RESULT 22

YK01_PYRHO
 ID YK01_PYRHO STANDARD; PRT; 147 AA.
 AC 057781;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hypothetical protein PH2001.
 GN PH2001.
 OS Pyrococcus horikoshii.
 OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococcaceae;
 OC Pyrococcus.
 OX NCBI_TaxID=53953;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98344137; PubMed=9679194;
 RA Kawanabayashi Y., Sawada M., Horikawa H., Haikawa Y., Hino Y.,
 RA Yamamoto S., Sakine M., Baba S.-I., Kosugi H., Hosoyama A., Nagai
 RA Sakai M., Ogura K., Otsuka R., Nakazawa H., Takamiya M., Ohfuku Y.
 RA Funahashi T., Tanaka T., Kudoh Y., Yamazaki J., Kushida N., Oguchi
 RA Aoki K.-I., Yoshizawa T., Nakamura Y., Robb F.T., Horikoshi K.,
 RA Masuchi Y., Shizuya H., Kikuchi H.;
 RT "Complete sequence and gene organization of the genome of a hyper
 thermophilic archaeobacterium, Pyrococcus horikoshii OT3.";
 RL DNA Res. 5:55-76(1998).

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 or send an email to license@isb-sib.ch).

 DR EMBL; AP000001; BAA31940.1;
 DR EMBL; AP000007; BAA31940.1; JOINED.
 DR EMBL; AP000007; BAA31943.1;
 DR EMBL; AP000001; BAA31943.1; JOINED.
 DR PIR; A71217; A71217.
 KW Hypothetical protein; Transmembrane; Complete proteome.
 FT TRANSMEM 41 61 POTENTIAL.
 FT TRANSMEM 67 87 POTENTIAL.
 SQ SEQUENCE 147 AA; 15324 MW; 247ED12FCE265B9 CRC64;

Query Match 2.5%; Score 7; DB 1; Length 147;

li O157:H7, and
eri.
obacteria; Gammaproteobacteria; Enterobacteriales;
ceae; Escherichia.
217992, 83334, 623;
N.A., AND SEQUENCE OF 21-40.
35; PubMed=2843433;
pe K.;
sequencing of the gene for the DNA-binding 17K protein
coli";
4 (1988).
N.A.
; STRAIN=K12 / MG1655;
17; PubMed=9278503;
Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,
ado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
is N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
; genome sequence of Escherichia coli K-12";
53-1474 (1997).
N.A.
; STRAIN=K12 / W3110;
ori H., Murayama N., Kataoka K., Yano M., Itoh T.,
nouchi H., Miki T., Hatada E., Fukuda R., Ichihara S.,
ino K., Nakata A., Yura T., Saueki G., Mizobuchi K.;
quencing of the Escherichia coli genome: analysis of the
(189,987 - 281,416bp) region";
1-1996) to the EMBL/GenBank/DBJ databases.
N.A.
;
ncan M., Allen E., Araujo R., Aparicio A., Chung E.,
rspei N., Hyman R., Kalman S., Komp C., Kurdi O.,
ew H., Lin D., Namath A., Oefner P., Roberts D.,
1-1996) to the EMBL/GenBank/DBJ databases.
N.A.
; STRAIN=O6:H1 / CFT073 / ATCC 700928;
34; PubMed=12471157;
urland V., Plunkett G. III, Redford P., Roesch P.,
les E.L., Liou S.-R., Boutin A., Hackett J., Stroud D.,
ose D.J., Zhou S., Schwartz D.C., Perna N.T.,
Donnenberg M.S., Blattner F.R.;
aic structure revealed by the complete genome sequence
ic Escherichia coli";
ad. Sci. U.S.A. 99:17020-17024 (2002).
N.A.
; STRAIN=O157:H7 / EDL933 / ATCC 700927;
35; PubMed=11206551;
unkett G. III, Burland V., Mau B., Glasner J.D.,
ew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,
kett J., Klink S., Boutin A., Shao Y., Miller L.,
Davis N.W., Lim A., Dimalanta E.T., Potamouais K.,
antharam T.S., Lin J., Yen G., Schwartz D.C.,
attner F.R.;
ce of enterohaemorrhagic Escherichia coli O157:H7";
533 (2001).
N.A.
; STRAIN=O157:H7 / RIMD 0509952;
31; PubMed=11258796;
kino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,
subo E., Nakayama K., Murata T., Tanaka M., Tobe T.,
i H., Honda T., Sasakawa C., Ogasawara N., Yasunaga T.,
ba T., Hattori M., Shingawa H.;
me sequence of enterohaemorrhagic Escherichia coli
nomic comparison with a laboratory strain K-12";

DNA Res. 8:11-22 (2001).
[8]
RL SEQUENCE OF 72-161 FROM N.A.
RN SPECIES=E.coli; STRAIN=K12 / MG1655;
RC MEDLINE=91100302; PubMed=1987124;
RX Dicker I.B., Seetharam S.R.;
RA "Cloning and nucleotide sequence of the fira gene and the fira200
RT allele from Escherichia coli";
RL J. Bacteriol. 173:334-344 (1991).
[9]
RN SEQUENCE OF 21-32.
RP SPECIES=E.coli; STRAIN=K12 / EMG2;
RC MEDLINE=97443975; PubMed=9298646;
RX Link A.J., Robison K., Church G.M.;
RA "Comparing the predicted and observed properties of proteins enco
RT in the genome of Escherichia coli K-12";
RL Electrophoresis 18:1259-1313 (1997).
[10]
RN SEQUENCE FROM N.A.
RP SPECIES=S.flexneri; STRAIN=301 / Serotype 2a;
RC MEDLINE=2272406; PubMed=12384590;
RA Jin Q., Yuan Z., Xu J., Wang Y., Shen Y., Lu W., Wang J., Liu H.,
Yang J., Yang F., Zhang X., Zhang J., Yang G., Wu H., Qu D., Dong
Sun L., Xue Y., Zhao A., Gao Y., Zhu J., Kan B., Ding K., Chen S.,
Cheng H., Yao Z., He B., Chen R., Ma D., Qiang B., Wen Y., Hou Y.,
Yu J.;
RA "Genome sequence of Shigella flexneri 2a: insights into pathogeni
RT through comparison with genomes of Escherichia coli K12 and O157.
RL Nucleic Acids Res. 30:4432-4441 (2002).
[11]
RN SEQUENCE FROM N.A.
RP SPECIES=S.flexneri; STRAIN=2457T / ATCC 700930 / Serotype 2a;
RC MEDLINE=22590274; PubMed=12704152;
RA Wei J., Goldberg M.B., Burland V., Venkatesan M.M., Deng W.,
Fournier G., Mayhew G.F., Plunkett G. III, Rose D.J., Darling A.,
Mau B., Perna N.T., Payne S.M., Runyen-Janecky L.J., Zhou S.,
Schwartz D.C., Blattner F.R.;
RA "Complete genome sequence and comparative genomics of Shigella
RT flexneri serotype 2a strain 2457T";
RL Infect. Immun. 71:2775-2786 (2003).
[12]
RN SIMILARITY TO S.TYPHIMURIUM OMPH.
RP SPECIES=E.coli;
RC MEDLINE=90201355; PubMed=2318304;
RA Hirvas L., Coleman J., Koski P., Vaara M.;
RT "Bacterial 'histone-like protein I' (HLP-I) is an outer membrane
constituent?";
RL FEBS Lett. 262:123-126 (1990).
CC -!- SUBUNIT: Homotetramer.
CC -!- SUBCELLULAR LOCATION: EITHER IN THE NUCLEOID (CHROMATIN) OR II
THE OUTER MEMBRANE.
CC -!- SIMILARITY: BELONGS TO THE OMPH/HLPa FAMILY.
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EMBL; M21118; AAA24630.1; -
DR EMBL; AE000127; AAC73289.1; -
DR EMBL; D83536; BAA77853.1; -
DR EMBL; U70214; AAB08607.1; -
DR EMBL; AE016755; AAN78707.1; -
DR EMBL; AF005194; AAG54480.1; -
DR EMBL; AF002550; BAB33603.1; -
DR EMBL; X54797; CAA38567.1; -
DR EMBL; X75465; CAA53207.1; -
DR EMBL; AE015054; AAN41830.1; -
DR EMBL; AE016978; AAP15711.1; -
DR FIR; D85502; D85502.

D90651.
DNEC17.
455; hlpA.
005632; Omph.
; Omph; 1.
Outer membrane; Signal; Complete proteome.
1 20
21 161 HISTONE-LIKE PROTEIN HLP-1.
1 AA; 17688 MW; 2A966BBD83F3E675 CRC64;
2.5%; Score 7; DB 1; Length 161;
arity 100.0%; Pred. No. 41;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ALA 68
|||
ALA 15
STANDARD; PRT; 170 AA.
[Rel. 28, Created]
[Rel. 28, Last sequence update]
[Rel. 41, Last annotation update]
protein D2007.4 in chromosome III.
s. elegans.
razos; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
Peizerinae; Caenorhabditis.
39;
[N.A.
[N2;
718; PubMed=7906398;
nscough R., Anderson K., Baynes C., Berks M.,
Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
Dear S., Du Z., Durbin R., Favello A., Fraser A.,
Jardner A., Green P., Hawkins T., Hillier L., Jier M.,
Jones M., Kershaw J., Kirsten J., Laister N.,
Lightning J., Lloyd C., Mortimore B., O'Callaghan M.,
Percy C., Rifken L., Roopra A., Saunders D., Showkhen R.,
Don N., Smith A., Smith M., Sonhammer E., Staden R.,
Hierry-Wieg J., Thomas K., Vaudin M., Vaughan K.,
Watson A., Weinstock L., Wilkinson-Sproat J.,
ntiguous nucleotide sequence from chromosome III of C.
38(1994).
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mail to license@isb-sib.ch).
AAA27999.1; -.
S44789.
7.4; CE00129.
protein.
70 AA; 19396 MW; 22301D7C65638135 CRC64;
2.5%; Score 7; DB 1; Length 170;
arity 100.0%; Pred. No. 43;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
JCL 227
|||
JCL 100

RESULT 27
LACB_BUBBU
ID LACB_BUBBU STANDARD; PRT; 180 AA.
AC P02755; 062822;
DT 21-JUL-1986 (Rel. 01, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Beta-lactoglobulin precursor (Beta-LG).
GN LGB.
OS Bubalus bubalis (Domestic water buffalo).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bubalus.
OX NCBI_TaxID=89462;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Mammary gland;
RX MEDLINE=99304500; PubMed=10376212;
RA Das P., Jain S., Nayak S., Apparao K.B.C., Totev S.M., Garg L.C.;
RT "Molecular Cloning and sequence analysis of the cDNA encoding
beta-lactoglobulin in Bubalus bubalis.";
RL DNA Seq. 10:105-108 (1999).
RN [2]
RP SEQUENCE OF 19-180.
RA Kolde H.-J., Liberatori J., Braunitzer G.;
RT "The amino acid sequence of the water buffalo beta-lactoglobulin
Milchweissenshaft 36:83-86 (1981).
RL
CC -!- FUNCTION: Primary component of whey, it binds retinol and is
probably involved in the transport of that molecule.
CC -!- SUBUNIT: Under physiological conditions beta-lactoglobulin e;
as an equilibrium mixture of monomeric and dimeric forms.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- TISSUE SPECIFICITY: Synthesized in mammary gland and secret
in milk.
CC -!- PTM: Alternate disulfide bonds occur in equal amounts.
CC -!- SIMILARITY: Belongs to the lipocalin family.
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EMBL; AJ005429; CAA06532.1; -.
DR InterPro; IPR002345; Lipocalin.
DR InterPro; IPR000566; Lipocalin_cytfabp.
DR Pfam; PF00061; lipocalin; 1.
DR PRINTS; PR00179; LIPOCALIN.
DR PROSITE; PS00213; LIPOCALIN; 1.
KW Milk; Whey; Retinol-binding; Transport; Lipocalin; Signal.
FT SIGNAL
FT CHAIN 1 180 BETA-LACTOGLOBULIN.
FT DISULFID 84 178
FT DISULFID 124 137
FT DISULFID 124 139
FT SEQUENCE 180 AA; 20023 MW; 6836C97B2C2E33CF CRC64;
ALTERNATE.
Query Match 2.5%; Score 7; DB 1; Length 180;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 7; Conservative 0; Mismatches 0; Indels 0;
QY 63 IGLALAC 69
DB 8 IGLALAC 14
RESULT 28
LACB_CAPHI
ID LACB_CAPHI STANDARD; PRT; 180 AA.
AC P02756;

rel. 01, Created)
 rel. 22, Last sequence update)
 rel. 43, Last annotation update)
 nulin precursor (Beta-LG).
 (Goat).
 azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 eria; Cetartiodactyla; Ruminantia; Pecora; Bovoidea;
 nae; Capra.
 5;
 N.A.
 agrus; TISSUE=Mammary gland;
 59; PubMed=8226387;
 all A., Sanchez A.;
 sequencing of the cDNA encoding goat beta-
 .;
 71:2832-2832(1993).
 N.A.
 Kim J., Yu M.;
 I-1993) to the EMBL/GenBank/DBJ databases.
 N.A.
 51; PubMed=7699130;
 A.; Sanchez A.;
 nence of the caprine beta-lactoglobulin gene";
 77:3493-3497(1994).
 1-180.
 11; PubMed=511095;
 unitzer G., Schrank B., Stangl A.;
 d sequence of goat beta-lactoglobulin";
 2. Physiol. Chem. 360:1595-1604(1979).
 Primary component of whey, it binds retinol and is
 involved in the transport of that molecule.
 Under physiological conditions beta-lactoglobulin exists
 librium mixture of monomeric and dimeric forms.
 AR LOCATION: Secreted.
 ICIFICITY: Synthesized in mammary gland and secreted
 nate disulfide bonds occur in equal amounts.
 : Belongs to the lipocalin family.
 NT entry is copyright. It is produced through a collaboration
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 il to license@isb-sib.ch).

 CAA41385.1; -
 CAA79623.1; -
 CAA79624.1; -
 CAA83946.1; -
 AGGT.
 1800.
 02345; Lipocalin.
 00566; Lipocalin_cytFABP.
 ; Lipocalin; 1.
 79; LIPOCALIN.
 213; LIPOCALIN; 1.
 etinol-binding; Transport; Lipocalin; Signal.
 1 18
 19 180 BETA-LACTOGLOBULIN.
 34 178
 24 137
 24 139 ALTERNATE.
) AA; 19975 MW; C2449BH02A1A80F1 CRC64;
 2.5%; Score 7; DB 1; Length 180;

Best Local Similarity 100.0%; Pred. No. 45;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G
 QY 63 LGLALAC 69
 Db 8 LGLALAC 14
 RESULT 29
 LACB SHEEP
 ID LACB SHEEP STANDARD; PRT; 180 AA.
 AC P02757;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 01-AUG-1988 (Rel. 08, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Beta-lactoglobulin 1/B, 2/A, and 3/C precursor.
 OS Ovis aries (Sheep), and
 OS Ovis orientalis musimon (Mouflon).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovoidea
 OC Bovidae; Caprinae; Ovis.
 OX NCBI_TaxID=9940, 9938;
 RN [1]
 RP SEQUENCE FROM N.A. (BLG 1 AND 2).
 RC SPECIES=Sheep;
 RX MEDLINE=88172489; PubMed=3351935;
 RA Ali S., Clark A.J.;
 RT "Characterization of the gene encoding ovine beta-lactoglobulin.
 RT Similarity to the genes for retinol binding protein and other
 RT secretory proteins.";
 RL J. Mol. Biol. 199:415-426(1988).
 RN [2]
 RP SEQUENCE FROM N.A. (BLG 1).
 RC SPECIES=Sheep;
 RX MEDLINE=87049827; PubMed=3096387;
 RA Gaye P., Hue-Delahaie D., Mercier J.-C., Soulier S., Vilotte J.-L
 RA Furet J.-P.;
 RT "Ovine beta-lactoglobulin messenger RNA: nucleotide sequence and
 RT levels during functional differentiation of the mammary gland.";
 RL Biochimie 68:1097-1107(1986).
 RN [3]
 RP SEQUENCE FROM N.A. (BLG 1).
 RC SPECIES=Sheep;
 RX MEDLINE=89057492; PubMed=3194215;
 RA Harris S., Ali S., Anderson S., Archibald A.L., Clark A.J.;
 RT "Complete nucleotide sequence of the genomic ovine beta-lactoglob
 RT gene.";
 RL Nucleic Acids Res. 16:10379-10380(1988).
 RN [4]
 RP SEQUENCE FROM N.A. (BLG 1 AND 2).
 RC SPECIES=Sheep;
 RX MEDLINE=91007276; PubMed=1976573;
 RA Ali S., McClenaghan M., Simons J.P., Clark A.J.;
 RT "Characterisation of the alleles encoding ovine beta-lactoglobuli
 RT and B.";
 RL Gene 91:201-207(1990).
 RN [5]
 RP SEQUENCE OF 19-180 (BLG 2).
 RC SPECIES=Sheep;
 RX MEDLINE=80219294; PubMed=6155855;
 RA Preaux G., Braunitzer G., Kolde H.-J.;
 RT "Primary structure of ovine beta-lactoglobulin.";
 RL Arch. Int. Physiol. Biochim. 88:845-846(1980).
 RN [6]
 RP SEQUENCE OF 19-180 (BLG 3).
 RC SPECIES=Sheep;
 RX MEDLINE=89374823; PubMed=2775495;
 RA Erhardt G., Godovac-Zimmermann J., Conti A.;
 RT "Isolation and complete primary sequence of a new ovine wild-type
 RT beta-lactoglobulin C.";
 RL Biol. Chem. Hoppe-Seyler 370:757-762(1989).
 RN [7]
 RP SEQUENCE OF 19-180 (BLG B).

```

musimon;
1996; PubMed=3426802;
rmann J., Conti A., Napolitano L.;
: amino-acid sequence of dimeric beta-lactoglobulin from
: ammon musimon) milk.;
: ioppe-Seyler 368:1313-1319(1987).
: LACTOGLOBULIN IS THE PRIMARY COMPONENT OF WHEY, IT
: INOL AND IS PROBABLY INVOLVED IN THE TRANSPORT OF
: ECULE.
Under physiological conditions beta-lactoglobulin exists
in equilibrium mixture of monomeric and dimeric forms.
Dimeric disulfide bonds occur in equal amounts.
: Belongs to the lipocalin family.
-----
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-profit institutions as long as its content is in no way
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: mail to license@isb-sib.ch).
-----
: CAA28204.1; -.
: CAA31305.1; -.
: CAA30059.1; ALT SEQ.
: CAA30059.1; JOINED.
: CAA30059.1; JOINED.
: CAA30059.1; JOINED.
: CAA30059.1; JOINED.
: CAA30059.1; JOINED.
: AAA31510.1; -.
: AAA31510.1; JOINED.
: AAA31510.1; JOINED.
: AAA31510.1; JOINED.
: LGSB.
: IBSQ.
R002345; Lipocalin.
R000566; Lipocalin_cytFABP.
1; Lipocalin; 1.
179; LIPOCALIN.
0213; LIPOCALIN; 1.
Retinol-binding; Transport; Signal; Lipocalin.
1 18
19 180 BETA-LACTOGLOBULIN.
84 178
124 137
124 139
38 166
166 166
30 AA; 19921 MW; BABCB2E89E757333 CRC64;
2.5%; Score 7; DB 1; Length 180;
larity 100.0%; Pred. No. 45;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ALAC 69
|||||
ALAC 14
STANDARD; PRT; 181 AA.
(Rel. 42, Created)
(Rel. 42, Last sequence update)
(Rel. 42, Last annotation update)
phoribosyltransferase (EC 2.4.2.7) (APRT).
2.
neidensis.
oteobacteria; Gammaproteobacteria; Alteromonadales;
ceae; Shewanella.
0863;

```

```

RN SEQUENCE FROM N.A.
RP STRAIN=MR-1;
RX MEDLINE=22397686; PubMed=12368813;
RA Heidelberg J.F., Paulsen I.F., Nelson K.E., Gaidos E.J., Nelson R.
RA Read T.D., Eisen J.A., Seshadri R., Ward N., Methe B., Clayton R.
RA Meyer T., Tsapin A., Scott J., Bearan M., Brinkac L., Daugherty
RA DeBoy R.T., Tadoin R.J., Durkin A.S., Haft D.H., Kolonay J.F.,
RA Madupu R., Peterson J.D., Unayam L.A., White O., Wolf A.M.,
RA Vamathevan J., Weidman J., Imbraim M., Lee K., Berry K., Lee C.,
RA Mueller J., Khouri H., Gill J., Uterback T.R., McDonald L.A., C.
RA Feldblyum T.V., Smith H.O., Venter J.C., Nealon K.H., Fraser C.
RT "Genome sequence of the dissimilatory metal ion-reducing bacteri
Shewanella oneidensis."
RT Nat. Biotechnol. 20:1118-1123(2002).
CC -!- FUNCTION: Catalyzes a salvage reaction resulting in the form
of AMP, that is energetically less costly than de novo synthe
CC -!- CATALYTIC ACTIVITY: AMP + diphosphate = adenine + 5-phospho-
D-ribose 1-diphosphate.
CC -!- PATHWAY: Purine salvage.
CC -!- SUBUNIT: Homodimer (By similarity).
CC -!- SUBCELLULAR LOCATION: Cytoplasmic.
CC -!- SIMILARITY: Belongs to the purine/pyrimidine
phosphoribosyltransferase family.
-----
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the European Bioinformatics Institute. There are no restrictio
use by non-profit institutions as long as its content is i
modified and this statement is not removed. Usage by and for
entities requires a license agreement (See http://www.isb-sib.ch
or send an email to license@isb-sib.ch).
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EMBL; AE015643; AAN55062.1; -.
TIGR; SO2012; -.
HAMAP; MF_00004; -.
DR InterPro; IPR005764; Ade phspho trans.
DR InterPro; IPR002375; Pr/py rp trans.
DR InterPro; IPR000836; Prtransferase.
DR Pfam; PF00156; Pribosyltran; 1.
DR TIGRFAMs; TIGR01090; apt; 1.
DR PROSITE; PS00103; PUR PYR PR TRANSFER; 1.
KW Transferase; Glycosyltransferase; Purine salvage; Complete prote
SQ SEQUENCE 181 AA; 19543 MW; C4255A59C4632CA4 CRC64;
Query Match 2.5%; Score 7; DB 1; Length 181;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 7; Conservative 0; Mismatches 0; Indels 0;
QY 58 PLALGIG 64
|||||
Db 71 PLALGIG 77
RESULT 31
DEF1 BIFLO
ID DEF1 BIFLO STANDARD; PRT; 217 AA.
AC Q8G534;
DT 15-MAR-2004 (Rel. 43, Created)
DT 15-MAR-2004 (Rel. 43, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Peptide deformylase 1 (EC 3.5.1.88) (PDF 1) (Polypeptide deformy
1).
DE DE
GN DEF1 OR BL1186.
OS Bifidobacterium longum.
OC Bacteria; Actinobacteria; Actinobacteridae; Bifidobacteriales;
OC Bifidobacteriaceae; Bifidobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=NCC 2705;
RX MEDLINE=22294977; PubMed=12381787;
RA Schell M.A., Karmirantzou M., Snel B., Vilanova D., Berger B.,

```

Allen M.-C., Desiere F., Bork P., Delley M.,
 Arigoni F.;
 sequence of Bifidobacterium longum reflects its adaptation
 to the gastrointestinal tract.";
 J. Biol. Chem. 274:14422-14427 (2002).
 Removes the formyl group from the N-terminal Met of
 the protein. Requires at least a dipeptide for an
 active site. N-terminal L-methionine is a
 rate of reaction. The enzyme has broad specificity at
 the for activity but the enzyme has broad specificity at
 itions (By similarity).
 ACTIVITY: Formyl-L-methionyl peptide + H₂O = formate +
 peptide.
 Binds 1 iron(II) ion (By similarity).
 Y: Belongs to the polypeptide deformylase family.
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 ail to license@isb-sib.ch.
 3; AAN24991.1; -;
 63; -; 1.
 000181; Fmet deformylase.
 ; Pep deformylase; 1.
 76; PDEFORMLASE;
 844; Pep deformylase; 1.
 nthesis; Hydrolase; Iron; Complete proteome.
 72 172 BY SIMILARITY.
 29 129 IRON (BY SIMILARITY).
 71 171 IRON (BY SIMILARITY).
 75 175 IRON (BY SIMILARITY).
 7 AA; 24443 MW; 52F5B469B6F74163 CRC64;
 2.5%; Score 7; DB 1; Length 217;
 arity 100.0%; Pred. No. 53;
 onservative 0; Mismatches 0; Indels 0; Gaps 0;
 ALA 68
 |||||
 ALA 86
 STANDARD; PRT; 230 AA.
 Rel. 41, Created)
 Rel. 41, Last sequence update)
 Rel. 42, Last annotation update)
 sport complex protein rnfE.
 54 OR STY1668 OR T1322.
 himirium, and
 phi.
 teobacteria; Gammaproteobacteria; Enterobacteriales;
 aceae; Salmonella.
 2, 601;
 N.A.
 himirium; STRAIN=LT2 / SGSC1412 / ATCC 700720;
 948; PubMed=11677609;
 , Sanderson K.E.; Spieth J., Clifton S.W., Latreille P.,
 Porwollik S., Ali J., Dante M., Du F., Hou S., Layman D.,
 Guyen C., Scott K., Holmes A., Grewal N., Mulvaney E.,
 H., Florea L., Miller W., Stoneking T., Nhan M.,
 Wilson R.K.;
 some sequence of Salmonella enterica serovar Typhimurium
 2-856(2001).
 I N.A.

SPECIES=S.typhi; STRAIN=CT18;
 MEDLINE=21534947; PubMed=11677608;
 RA Parkhill J., Dougan G., James K.D., Thomson N.R., Pickard D., Wai
 RA Churcher C., Mungall K.L., Bentley S.D., Holden M.T.G., Sebaihia
 RA Baker S., Baeham D., Brooks K., Chillingworth T., Connor P.,
 RA Cronin A., Davis P., Davies R.M., Dowd L., White N., Farrar J.,
 RA Feltwell T., Hamlin N., Haque A., Hien T.T., Holroyd S., Jagels K
 RA Krogh A., Larsen T.S., Leather S., Moule S., O'Gaora P., Parry C.
 RA Quail M.A., Rutherford K., Simmonds M., Skelton J., Stevens K.,
 RA Whitehead S., Barrall B.G.;
 RT "Complete genome sequence of a multiple drug resistant Salmonella
 RT enterica serovar Typhi CT18.";
 RL Nature 413:848-852(2001).
 RP [3]
 RP SEQUENCE FROM N.A.
 RP SPECIES=S.typhi; STRAIN=Ty2 / ATCC 700931;
 RX MEDLINE=22531367; PubMed=12644504;
 RA Deng W., Liou S.-R., Plunkett G. III, Mayhew G.F., Rose D.J.,
 RA Burland V., Kodyanni V., Schwartz D.C., Blattner F.R.;
 RT "Comparative genomics of Salmonella enterica serovar Typhi strain
 RT and CT18";
 RL J. Bacteriol. 185:2330-2337(2003).
 CC -!- FUNCTION: May be part of a membrane complex involved in elect
 CC transport (By similarity).
 CC -!- SUBUNIT: Composed of at least six subunits; rnfA, rnfB, rnfC,
 CC rnfD, rnfE and rnfG (By similarity).
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Inner membra
 CC (Potential).
 CC -!- SIMILARITY: Belongs to the nqrDE/rnfAE family.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a coll
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 CC the European Bioinformatics Institute. There are no restriction
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 CC entities requires a license agreement (See <http://www.isb-sib.ch/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; AE008763; AAL20376.1; -;
 DR EMBL; AL627271; CAD01913.1; -;
 DR EMBL; AE016838; AAC68972.1; -;
 DR StyGene; SG????; rnfE.
 DR HAMAP; MF 00478; -; 1.
 DR InterPro; IPR003667; Rnf_Nqr.
 DR Pfam; PF02508; Rnf_Nqr; I
 KW Electron transport; Transmembrane; Inner membrane; Complete prote
 FT TRANSMEM 34 56 POTENTIAL.
 FT TRANSMEM 69 87 POTENTIAL.
 FT TRANSMEM 91 113 POTENTIAL.
 FT TRANSMEM 126 148 POTENTIAL.
 FT TRANSMEM 183 205 POTENTIAL.
 SQ SEQUENCE 230 AA; 24318 MW; E198B4CEA13F249E CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 230;
 Best Local Similarity 100.0%; Pred. No. 56;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; C
 QY 60 ALGGLA 66
 |||||
 DB 38 ALGGLA 44
 RESULT 33
 RNFE_ECO57
 ID RNFE_ECO57 STANDARD; PRT; 231 AA.
 AC P58344;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Electron transport complex protein rnfE.
 GN RNFE OR Z2642 OR ECS2341.
 OS Escherichia coli O157:H7.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;

aceae; Escherichia.

3334;

4 N.A.

17 / EBL933 / ATCC 700927;

1935; PubMed:11206551;

Plunkett G. III, Burland V., Mau B., Glasner J.D.,

lyhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,

ickett J., Klink S., Boutin A., Shao Y., Miller L.,

.. Davis N.W., Lim A., Dimalanta E.T., Potamoumis K.,

Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,

Blattner F.R.;

ence of enterohaemorrhagic Escherichia coli O157:H7.;

29-533 (2001).

4 N.A.

17 / RMD 0509952;

5231; PubMed:11258796;

akino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,

itubo E., Nakayama K., Murata T., Tanaka M., Tobe T.,

mi H., Honda T., Sasakawa C., Ogasawara N., Yasunaga T.,

iba T., Hattori M., Shinagawa H.;

ome sequence of enterohemorrhagic Escherichia coli

genomic comparison with a laboratory strain K-12.;

[-22 (2001).

: May be part of a membrane complex involved in electron

: (By similarity).

Composed of at least six subunits; rnfA, rnfB, rnfC,

IE and rnfG (By similarity).

AR LOCATION: Integral membrane protein. Inner membrane

IV).

IV): Belongs to the nqrDE/rnfAE family.

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36; AAG56621.1; -.

58; BAB35764.1; -.

A85770.

E90921.

178; -; 1.

3003667; Rnf Nqr.

3; Rnf-Nqr; 1.

isport; Transmembrane; Inner membrane; Complete proteome.

1 38 PERIPLASMIC (POTENTIAL).

39 59 POTENTIAL.

60 62 CYTOPLASMIC (POTENTIAL).

63 83 POTENTIAL.

84 85 PERIPLASMIC (POTENTIAL).

86 106 POTENTIAL.

107 124 CYTOPLASMIC (POTENTIAL).

125 145 POTENTIAL.

146 181 PERIPLASMIC (POTENTIAL).

182 202 POTENTIAL.

203 231 CYTOPLASMIC (POTENTIAL).

31 AA; 24489 MW; D4A2CA2D292604C3 CRC64;

2.5%; Score 7; DB 1; Length 231;

larity 100.0%; Pred. No. 56;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LGLA 66

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LGLA 44

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4 85 PERIPLASMIC (POTENTIAL).
6 106 POTENTIAL.
7 124 CYTOPLASMIC (POTENTIAL).
5 145 POTENTIAL.
6 181 PERIPLASMIC (POTENTIAL).
2 202 POTENTIAL.
3 231 CYTOPLASMIC (POTENTIAL).
AA; 24459 MW; CFA37A2D292604C3 CRC64;
2.5%; Score 7; DB 1; Length 231;
urity 100.0%; Pred.No.56;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
ALA 66
||
ALA 44
STANDARD; PRT; 233 AA.
rel. 34, Created)
rel. 42, Last sequence update)
rel. 42, Last annotation update)
III (EC 3.1.26.3) (RNase III).
btti.
eobacteria; Gammaproteobacteria; Legionellales;
Coxiella.
;
N.A.
/;
751; PubMed=7830573;
fer T.A., Powell B.S., Court D.L.;
the rnc locus of Coxiella burnetii.";
l. 14:291-300(1994).
N.A.
ile phase I / RSA 493;
557; PubMed=12704232;
Paulsen I.T., Eisen J.A., Read T.D., Nelson K.E.,
Vard N.L., Tettelin H., Daviden T.M., Beanan M.J.,
ugherty S.C., Brinkac L.M., Madupu R., Dodson R.J.,
Lee K.H., Carty H.A., Scanlan D., Heinzen R.A.,
Samuel J.E., Fraser C.M., Heidelberg J.F.;
ome sequence of the Q-fever pathogen, Coxiella
cad. Sci. U.S.A. 100:5455-5460(2003).
Digests double-stranded RNA. Involved in the processing
nal RNA precursors and of some mRNAs (By similarity).
ACTIVITY: Endonucleolytic cleavage to 5'-
noster.
AR LOCATION: Cytoplasmic.
Y: Contains 1 DBM (double-stranded RNA-binding) domain.
Y: Contains 1 RNase III domain.
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AAA69690.1; -;
4: AA091000.1; -;
S60767.
;-;
04; -; 1.
001159; DS RBD.
000999; RNase_III.

DR Pfam; PF00035; dsrm; 1.
DR Pfam; PF00636; Ribonuclease_3; 1.
DR SMART; SM00358; DSRM; 1.
DR SMART; SM00535; RIBOC; 1.
DR PROSITE; PS0137; DS RBD; 1.
DR PROSITE; PS00517; RNase_3; 1.
DR PROSITE; PS0142; RNase_3; 2; 1.
KW Hydrolase; Nuclease; Endonuclease; RNA-binding; Complete proteome
FT DOMAIN 4 126 RNase III.
FT DOMAIN 204 220 DBM.
FT CONFLICT 116 116 A -> T (IN REF. 1).
SQ SEQUENCE 233 AA; 26199 MW; 1A11CB5FD960784F CRC64;
Query Match 2.5%; Score 7; DB 1; Length 233;
Best Local Similarity 100.0%; Pred.No.56;
Matches 7; Conservative 0; Mismatches 0; Indels 0; G
QY 10 ARRLPLP 16
|||
Db 164 ARRLPLP 170
RESULT 36
RNFE YERPE STANDARD; PRT; 233 AA.
AC Q8ZED4;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Electron transport complex protein rnfE.
GN RNFE OR YPO2240 OR Y2081.
OS Versinia pestis.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Versinia.
OX NCBI_TaxID=632;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CO-92 / Biovar Orientalis;
RX MEDLINE=21470413; PubMed=11586360;
RA Parkhill J., Wren B.W., Thomson N.R., Titball R.W., Holden M.T.G.
RA Prentice M.B., Sebaihia M., James K.D., Churcher C., Mungall K.L.
RA Baker S., Basham D., Bentley S.D., Brooks K., Cerdeno-Tarraga A.M.
RA Chillingworth T., Cronin A., Davies R.M., Davis P., Dougan G.,
RA Feltwell T., Hamlin N., Holroyd S., Jagels K., Karlyshev A.V.,
RA Leather S., Moule S., Oyston P.C.F., Quail M.A., Rutherford K.,
RA Simmonds M., Skelton J., Stevens K., Whitehead S., Barrell B.G.;
RT "Genome sequence of Versinia pestis, the causative agent of plagu
RL Nature 413:523-527(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=KIM5 / Biovar Mediaevalis;
RX MEDLINE=22137863; PubMed=12142430;
RA Deng W., Burland V., Plunkett G. III, Boutin A., N
RA Perna N.T., Rose D.J., Mau B., Zhou S., Schwartz /
RA Fetherston J.D., Lindler L.E., Brubaker R.R., Pls
RA Straley S.C., McDonough K.A., Nilles M.L., Matso'
RA Perry R.D.;
RT "Genome sequence of Versinia pestis KIM.";
RL J. Bacteriol. 184:4601-4611(2002).
CC -!- FUNCTION: May be part of a membrane comple
transport (By similarity).
CC -!- SUBUNIT: Composed of at least six subunit
rnfD, rnfE and rnfG (By similarity).
CC -!- SUBCELLULAR LOCATION: Integral membrane
(Potential).
CC -!- SIMILARITY: Belongs to the nqrDE/rnfAE /
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mail to license@isb-sib.ch).

51; CAC91046.1; -;
 10; AAM85645.1; -;
 AB0273.
 478; -; 1.
 R003667; Rnf Nqr.
 9; Rnf-Nqr; 1.
 asport; Transmembrane; Inner membrane; Complete proteome.
 34 56 POTENTIAL.
 69 87 POTENTIAL.
 91 113 POTENTIAL.
 126 148 POTENTIAL.
 184 206 POTENTIAL.
 33 AA; 24587 MW; 491E18F335B8CB90 CRC64;

2.5%; Score 7; DB 1; Length 233;
 larity 100.0%; Pred. No. 56; 0; Indels 0; Gaps 0;
 Conservative 0; Mismatches 0;

LGLA 66
 ||||
 LGLA 44

46; STANDARD; PRT; 235 AA.

(Rel. 35, Created)
 (Rel. 35, Last sequence update)
 (Rel. 41, Last annotation update)
 nsport complex protein rnfE.

88.
 influenzae.
 oteobacteria; Gammaproteobacteria; Pasteurellales;
 eae; Haemophilus.

27;

M N.A.

KW20 / ATCC 51907;
 0630; PubMed=7542800;
 R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
 R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
 Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
 Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
 Phillips C.A., Spriggs T., Hedblom E., Corton M.D.,
 R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
 Ritzman J.L., Fuhrmann J.L., Geoghegan N.S.M.,
 McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,

e random sequencing and assembly of Haemophilus influenzae

496-512(1995).

t May be part of a membrane complex involved in electron

t (By similarity).

Composed of at least six subunits; rnfA, rnfB, rnfC,

FE and rnfG (By similarity).

LAR LOCATION: Integral membrane protein. Inner membrane

al).

TY: Belongs to the nqrDE/rnfAE family.

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; AAC23334.1; -;

I64174.

;

DR HAMAP; MF_00478; -; 1.
 DR InterPro; IPR003667; Rnf Nqr.
 DR Pfam; PF02508; Rnf-Nqr; 1.
 KW Electron transport; Transmembrane; Inner membrane; Complete prot
 FT TRANSMEM 63 83 POTENTIAL.
 FT TRANSMEM 93 113 POTENTIAL.
 FT TRANSMEM 117 137 POTENTIAL.
 FT TRANSMEM 152 172 POTENTIAL.
 FT TRANSMEM 206 226 POTENTIAL.
 SQ SEQUENCE 235 AA; 25845 MW; C054FE596647837A CRC64;

Query Match 2.5%; Score 7; DB 1; Length 235;
 Best Local Similarity 100.0%; Pred. No. 57;
 Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 60 ALGLGLA 66

Db 62 ALGLGLA 68

RESULT 38

TN14 MOUSE STANDARD; PRT; 239 AA.

ID TN14 MOUSE STANDARD; PRT; 239 AA.
 AC QSOYH9;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Tumor necrosis factor ligand superfamily member 14.
 GN TNFSF14 OR LIGHT.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mu
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20165223; PubMed=10700230;
 RA Tamada K., Shimozaaki K., Chapoval A.I., Zhu G., Sica G., Flies I
 RA Boone T., Hsu H., Fu Y.-X., Nagata S., Ni J., Chen L.;
 RT "Modulation of T-cell-mediated immunity in tumor and graft-versu
 RT disease models through the LIGHT co-stimulatory pathway.";
 RL Nat. Med. 6:283-289(2000).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX TISSUE=Fetal liver;
 RC MEDLINE=20354998; PubMed=10894944;
 RA Misawa K., Nosaka T., Kojima T., Hirai M., Kitamura T.;
 RT "Molecular cloning and characterization of a mouse homolog of hu
 RT TNFSF14, a member of the TNF superfamily.";
 RL Cytogenet. Cell Genet. 89:89-91(2000).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX TISSUE=Lymphoma;
 RC Force W.R., Todd P.K., Mikayama T.;
 RT "Mouse LIGHT; molecular genetics, ligand binding and expression.
 RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.

CC -1- FUNCTION: Cytokine that binds to TNFRSF3/LTBR. Binding to th
 CC decoy receptor TNFRSF6B modulates its effects. Activates NFK
 CC and stimulates the proliferation of T cells.
 CC -1- SUBUNIT: Homotrimer (By similarity).
 CC -1- SUBCELLULAR LOCATION: Type II membrane protein and secreted
 CC similarity).
 CC -1- PTM: The soluble form derives from the membrane form by
 CC proteolytic processing.
 CC -1- SIMILARITY: Belongs to the tumor necrosis factor family.
 CC
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5; AAF76453.1; -;
 5; BAA8559.1; -;
 3; AAF36722.1; -;
 4TSV.
 317; Tnfstf14.
 006053; TNF abc.
 006052; TNF_family.
 009883; TNF_like.
 003636; TNF_subf.
 ; TNF; 1.
 34; TNECROSISFCT.
 012; TNF_subf; 1.
 7; TNF; 1.
 251; TNF_1; FALSE_NEG.
 049; TNF_2; 1.
 asmembrane; Glycoprotein; Signal-anchor.
 1 239 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
 MEMBER 14, MEMBRANE FORM.
 32 239 TUMOR NECROSIS FACTOR LIGAND SUPERFAMILY
 MEMBER 14, SOLUBLE FORM.
 1 37 CYTOPLASMIC (POTENTIAL).
 38 58 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
 (POTENTIAL).
 59 239 EXTRACELLULAR (POTENTIAL).
 31 82 CLEAVAGE (POTENTIAL).
 52 187 POTENTIAL.
 30 100 N-LINKED (GLCNAC. . .) (POTENTIAL).
 31 191 N-LINKED (GLCNAC. . .) (POTENTIAL).
 3 AA; 26338 MW; 217874AC71AD6BE3 CRC64;
 2.5%; Score 7; DB 1; Length 239;
 arity 100.0%; Pred.No. 58;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 RPR 127
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 RPR 227
 STANDARD; PRT; 240 AA.
 Rel. 41, Created)
 Rel. 41, Last sequence update)
 Rel. 41, Last annotation update)
 sport complex protein rnfE.
 4.
 aruginosa.
 ceobacteria; Gammaproteobacteria; Pseudomonadales;
 eae; Pseudomonas.
 7;
 N.A.
 5692 / PA01;
 337; PubMed=10984043;
 Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warriner P.,
 Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
 Soltry L., Tolentino E., Westbrock-Wadman S., Yuan Y.,
 Culter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
 pence D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
 ler M.H., Hancock R.E.W., Lory S., Olson M.V.;
 ome sequence of *Pseudomonas aeruginosa* PA01, an
 pathogen.";
 9-964(2000).
 May be part of a membrane complex involved in electron
 (By similarity).
 Composed of at least six subunits; rnfA, rnfB, rnfC,
 E and rnfG (By similarity).
 AR LOCATION: Integral membrane protein. Inner membrane
 1).
 Y: Belongs to the nqrDE/rnfAE family.

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 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; AE004770; AAG06882.1; -;
 DR PIR; G83208; G83208.
 DR HAMAP; MF 00478; -; 1.
 DR InterPro; IPR003667; Rnf_Nqr.
 DR Pfam; PF02508; Rnf-Nqr; 1.
 KW Electron transport; Transmembrane; Inner membrane; Complete prote
 FT TRANSMEM 41 61 POTENTIAL.
 FT TRANSMEM 71 91 POTENTIAL.
 FT TRANSMEM 95 115 POTENTIAL.
 FT TRANSMEM 130 150 POTENTIAL.
 FT TRANSMEM 184 204 POTENTIAL.
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 Query Match 2.5%; Score 7; DB 1; Length 240;
 Best Local Similarity 100.0%; Pred.No. 58;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G
 Qy 60 ALGLGLA 66
 Db 40 ALGLGLA 46
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 |||||
 RESULT 40
 MOEB HAEIN
 ID MOEB HAEIN STANDARD; PRT; 243 AA.
 AC P45211;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Molybdopterin biosynthesis protein moeb.
 GN MOEB OR CHLN OR H11449.
 OS Haemophilus influenzae.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
 OC Pasteurellaceae; Haemophilus.
 OC NCBI_TaxID=727;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Rd / KW20 / ATCC 51907;
 RX MEDLINE=95350630; PubMed=7542800;
 RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.
 RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.
 RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
 RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
 RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
 RA Utterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C
 RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
 RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
 RA Venter J.C.;
 RT "Whole-genome random sequencing and assembly of *Haemophilus influ*
 RT Rd.";
 RL Science 269:496-512(1995).
 CC -!- FUNCTION: INVOLVED IN BIOSYNTHESIS OF A DEMOLYBDO COFACTOR
 CC (MOLYBDOPTERIN), NECESSARY FOR MOLYBDENZYMES. PLAYS A ROLE I
 CC ACTIVATION OF THE SMALL SUBUNIT OF THE MOLYBDOPTERIN CONVERTI
 CC FACTOR (MOAD) (BY SIMILARITY).
 CC -!- PATHWAY: Molybdenum cofactor biosynthesis.
 CC -!- SIMILARITY: BELONGS TO THE HESA/MOEB/THIF FAMILY.
 CC -----
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AAC23099.1; -.
C64124.
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3009036; MoeB.
3007901; MoeZ MoeB.
3000205; NAD.BS.
3000594; Thif_domain.
3; MoeZ_MoeB; 1.
3; Thif; 1.
Factor biosynthesis; Complete proteome.
13 AA; 26996 MW; 218A3382A975BDBD CRC64;
2.5%; Score 7; DB 1; Length 243;
larity 100.0%; Pred.No.59;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
NRQI 185
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NRQI 16
STANDARD; PRT; 244 AA.
70; Q99761;
(Rel. 29, Created)
(Rel. 43, Last annotation update)
beta (IT-beta) (Tumor necrosis factor C) (TNF-C) (Tumor
cor ligand superfamily member 3).
3 OR TNFC.
(Human).
stazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Primates; Catarrhini; Hominidae; Homo.
506;
M N.A. (ISOFORM 1), AND PARTIAL SEQUENCE.
1;
3881; PubMed-7916655;
.. Ngan-Ek A., Lawton P., Demarinis J., Tizard R.,
ession C., O'Brine-Greco B., Foley S.F., Ware C.F.;
beta, a novel member of the TNF family that forms a
complex with lymphotoxin on the cell surface."
356(1993).
M N.A. (ISOFORMS 1 AND 2).
5965; PubMed-9299492;
Renard N., Charlot C., Bienvenu J., Coiffier B.,
ion of two lymphotoxin beta isoforms expressed in human
1 lines and non-Hodgkin's lymphomas."
phys. Res. Commun. 238:273-276(1997).
M N.A. (ISOFORM 1).
, Milner C.M., Campbell R.D.;
r of the immunoglobulin superfamily and a V-ATPase G
amongst the predicted products of novel genes close to the
the human MHC."
EP-1997) to the EMBL/GenBank/DBJ databases.
M N.A. (ISOFORM 1).
dan A., Qin S., Shaffer T., James R., Ratcliffe A.,
ickhoff R., Loretz C., Madan A., Dors M., Young J.,
od L.;
the human major histocompatibility complex class III
CT-1999) to the EMBL/GenBank/DBJ databases.
M N.A. (ISOFORM 1).
amiya G., Oka A., Inoko H.;
s 2,229,817bp genomic DNA of 6p21.3 HLA class I region."
EP-1999) to the EMBL/GenBank/DBJ databases.

```

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[6]
SEQUENCE FROM N.A. (ISOFORMS 1 AND 2), AND VARIANTS GLU-70 AND
PRO-111.
RA Rieder M.J., Armel T.Z., Carrington D.P., Chung M.-W., Lee K.L.,
RA Poel C.L., Toth E.J., Yi Q., Nickerson D.A.;
RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.
[7]
SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS ARG-84 AND PHE-87.
RA Rieder M.J., Livingston R.J., Daniels M.R., Montoya M.A., Chung
RA Miyamoto K.E., Nguyen C.P., Nguyen D.A., Poel C.L., Robertson P.
RA Schackwitz W.S., Sherwood J.K., Wittrak L.A., Nickerson D.A.;
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Cytokine that binds to LTBR/TNFRSF3. May play a sp
CC role in immune response regulation. Provides the membrane an
CC for the attachment of the heterotrimeric complex to the cell
CC surface. Isoform 2 is probably non-functional.
CC -!- SUBUNIT: Heterotrimer of either two LTB and one LTA subunits
CC (less prevalent) one LTB and two LTA subunits.
CC -!- SUBCELLULAR LOCATION: Type II membrane protein (Potential).
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1;
CC IsoId=Q06643-1; Sequence=Displayed;
CC Name=2;
CC IsoId=Q06643-2; Sequence=VSP_006441, VSP_006442;
CC -!- TISSUE SPECIFICITY: Spleen and thymus.
CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
CC
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CC between the Swiss Institute of Bioinformatics and the EMBL ou
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EMBL; L11016; AAA99888.1; -
EMBL; U89922; AAC51769.1; -
EMBL; U79029; AAB37342.1; -
EMBL; L11015; AAA36191.1; -
EMBL; Y14768; CAA75069.1; -
EMBL; AF129756; RAD18089.1; -
EMBL; AP000505; BAB63395.1; -
EMBL; AY070219; AAL49954.1; -
EMBL; AY070219; AAL49955.1; -
EMBL; AY216497; AAO21134.1; -
PIR; A46066; A46066.
PIR; JCS645; JCS645.
HSSP; P01374; 1TNF.
Genew; HGNC:6711; LTB.
MIM; 600978; -.
GO; GO:0005102; F:receptor binding; TAS.
GO; GO:0015070; F:toxin activity; NAS.
GO; GO:0007267; P:cell-cell signaling; TAS.
GO; GO:0007165; P:signal transduction; TAS.
InterPro; IPR006053; TNF_abc.
InterPro; IPR006052; TNF_family.
InterPro; IPR008983; TNF_like.
InterPro; IPR003636; TNF_subf.
Pfam; PF00229; TNF; 1.
PRINTS; PR01234; TNECROSISFCT.
ProDom; PD002012; TNF_subf; 1.
SMART; SM00207; TNF; 1.
PROSITE; PS00251; TNF_1; 1.
PROSITE; PS00049; TNF_2; 1.
KW Cytokine; Transmembrane; Glycoprotein; Signal-anchor;
KW Alternative splicing; Polymorphism.
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FT TRANSMEM 19 48
FT SIGNAL-ANCHOR (TYPE-II MEMBRANE PRO
FT (POTENTIAL).
FT DOMAIN 49 244
FT EXTRACELLULAR (POTENTIAL).
FT CARBOHYD 222 222
FT VARSPPLIC 53 77
FT GLVTETADPGAQAQQGLGFKLPEE -> GLGFRS
(POTENTIAL).
N-LINKED (GLCNAC. .) (POTENTIAL).
GLVTETADPGAQAQQGLGFKLPEE -> GLGFRS
(POTENTIAL).

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 Missing (in isoform 2).
 /FTid=VSP_006442.
 G -> E.
 S -> R.
 /FTid=VAR_013025.
 /FTid=VAR_016331.
 L -> F.
 /FTid=VAR_016332.
 A -> P.
 /FTid=VAR_013026.
 DPGAQOQGL -> GLSAPSGRT (IN REF. 2;
 AAB37342).
 AA; 25390 MW; F41569459830ED4C CRC64;
 2.5%; Score 7; DB 1; Length 244;
 100.0%; Pred. No. 59;
 0; Mismatches 0; Indels 0; Gaps 0;
 YC 200
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 YC 137
 STANDARD; PRT; 244 AA.
 rel. 43, Created)
 rel. 43, Last sequence update)
 rel. 43, Last annotation update)
 eta (LT-beta) (Tumor necrosis factor C) (TNF-C) (Tumor
 or TNFC.
 as (Chimpanzee).
 asoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 peria; Primates; Catarrhini; Hominidae; Pan.
 18;
 N.A.
 102; PubMed=12493009;
 Shihina T., Anzai T., Kohara S., Inoko H.;
 genomic analysis of the MHC: the evolution of class I
 locks, diversity and complexity from shark to man.";
 190:95-122(2002).
 N.A.
 34; PubMed=12799463;
 na T., Kimura N., Yanagiya K., Kohara S., Shigenari A.,
 Galski J.K., Naruse T.K., Fujimori Y., Fukuzumi Y.,
 ashiro H., Iwanoto C., Umebara Y., Imanishi T.,
 K., Gojobori T., Bahram S., Inoko H.;
 sequencing of human and chimpanzee MHC class I regions
 ions/deletions as the major path to genomic
 ad. Sci. U.S.A. 100:7708-7713(2003).
 Cytokine that binds to LTR/TNFRSF3. May play a specific
 immune response regulation. Provides the membrane anchor
 attachment of the heterotrimeric complex to the cell
 by similarity).
 heterotrimer of either two LTB and one LTA subunits or
 valent) two LTA and one LTB subunits (BY similarity).
 AR LOCATION: Type II membrane protein (Potential).
 y: Belongs to the tumor necrosis factor family.
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CC EMBL; AB054536; BAB83881.1; -.
 DR EMBL; AB100082; BAC78156.1; -.
 DR InterPro; IPR006053; TNF_abc.
 DR InterPro; IPR006052; TNF_family.
 DR InterPro; IPR008983; TNF_like.
 DR InterPro; IPR003636; TNF_subf.
 DR Pfam; PF00229; TNF; 1.
 DR PRINTS; PR01234; TNECROSISFCT.
 DR ProDom; PD002012; TNF_subf; 1.
 DR SMART; SM00207; TNF; 1.
 DR PROSITE; PS00251; TNF_1; 1.
 DR PROSITE; PS0049; TNF_2; 1.
 KW Cytokine, transmembrane; Glycoprotein; Signal-anchor.
 FT CYTOPLASMIC (POTENTIAL).
 FT SIGNAL-ANCHOR (TYPE-II MEMBRANE PROT
 (POTENTIAL).
 FT DOMAIN 1 18
 FT TRANSMEM 19 48
 FT DOMAIN 49 244
 FT CARBOHYD 222 222
 FT N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 244 AA; 25420 MW; A4047858335D5B97 CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 244;
 Best Local Similarity 100.0%; Pred. No. 59;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G.
 QY 194 GLYLYYC 200
 DB 131 GLYLYYC 137
 |||||
 |||||
 RESULT 43
 LFTX XANCP STANDARD; PRT; 249 AA.
 ID LFTX XANCP STANDARD; PRT; 249 AA.
 AC QBP956; (Rel. 42, Created)
 DT 10-OCT-2003 (Rel. 42, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Leucyl-phenylalanyl-tRNA--protein transferase (EC 2.3.2.6) (L/F-
 DE transferase) (Leucyltransferase) (Phenylalanyltransferase).
 GN AAT OR XCC1969.
 OS Xanthomonas campestris (pv. campestris).
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
 CC Xanthomonadaceae; Xanthomonas.
 CX NCBI_TaxID=340;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 33913 / NCPPB 528;
 RC MEDLINE=22022145; PubMed=12024217;
 RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.
 RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.
 RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
 RA Camarotte G., Cannavan F., Cardozo J., Chambergo F., Ciapina L.P.
 RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorry H.
 RA Paria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
 RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
 RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.
 RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
 RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
 RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.,
 RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
 RA Spinola L.A.F., Takita M.A., Tamura R.B., Teixeira E.C., Tezza R.
 RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
 RA Stubal J.C., Kitajima J.P.;
 RT "Comparison of the genomes of two Xanthomonas pathogens with diff
 RT host specificities.";
 RL Nature 417:459-463(2002).
 CC -!- FUNCTION: Functions in the N-end rule pathway of protein
 CC degradation where it conjugates Leu, Phe and, less efficientl
 CC Met from aminoacyl-tRNAs to the N-termini of proteins contain
 CC an N-terminal arginine or lysine (By similarity).
 CC -!- CATALYTIC ACTIVITY: L-leucyl-tRNA + protein = tRNA + L-leucyl
 CC protein.
 CC -!- CATALYTIC ACTIVITY: L-phenylalanyl-tRNA + protein = tRNA + L-

OF THE SMALL SUBUNIT OF THE MOLYBDOPTEIN CONVERTING
 (MAD).
 Polymorphism cofactor biosynthesis.
 BELONGS TO THE HESA/MOE/THIF FAMILY.
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 ail to license@isb-sib.ch).

 AAA96530.1; --
 5; AAL19781.1; --
 565; moeb.
 009036; Moeb.
 007901; Moeb Moeb.
 00205; NAD BS.
 00594; Thif domain.
 ; Moeb Moeb; 1.
 ; Thif; 1.
 Factor biosynthesis; Complete proteome.
 9 9 M -> I (IN REF. 1).
 38 38 G -> R (IN REF. 1).
 L7 117 S -> A (IN REF. 1).
 59 169 N -> T (IN REF. 1).
 21 221 G -> E (IN REF. 1).
 ; AA; 26903 MW; 0F0050831D537AD2 CRC64;
 2.5%; Score 7; DB 1; Length 249;
 arity 100.0%; Pred.No. 60;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 QI 185
 ||||
 QI 16
 STANDARD; PRT; 257 AA.
 rel. 39, Created)
 rel. 39, Last sequence update)
 rel. 41, Last annotation update)
 aride core biosynthesis glycosyl transferase kdtx

 escents.
 eobacteria; Gammaproteobacteria; Enterobacteriales;
 aceae; Serratia.
 5;
 N.A.
 003; PubMed=8824620;
 pique N.; Climent N., Ferrer S., Merino S., Rubires X.,
 ague M.;
 Characterization of two Serratia marcescens genes
 core lipopolysaccharide biosynthesis.";
 178:5741-5747(1996).
 Lipopolysaccharide core biosynthesis.
 Y: BELONGS TO THE GLYCOSYLTRANSFERASE FAMILY 2. WAAE/KDTX

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 res a license agreement (See <http://www.isb-sib.ch/announce/>
 ail to license@isb-sib.ch).

CC EMBL; US2844; AAC44433.1; --
 DR InterPro; IPR001173; Glyco trans 2.
 DR Pfam; PF00535; Glycos transf 2; 1.
 DR Lipopolysaccharide biosynthesis; Transferase; Glycosyltransferase
 KW SEQUENCE 257 AA; 29233 MW; D40D7B57E002F990 CRC84;
 SQ
 Query Match 2.5%; Score 7; DB 1; Length 257;
 Best Local Similarity 100.0%; Pred.No. 62;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G
 QY 71 GLLAVV 77
 |||||
 Db 229 GLLAVV 235
 RESULT 47
 CN09 HUMAN STANDARD; PRT; 277 AA.
 ID CN09 HUMAN Q86U09; Q8WUC0; Q9BU67; Q9NSU8;
 AC Q86T03; Q86U09; Q8WUC0; Q9BU67; Q9NSU8;
 DT 10-OCT-2003 (Rel. 42, Created)
 DT 10-OCT-2003 (Rel. 42, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Hypothetical protein Cl4orf9.
 GN Cl4ORF9.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
 RC TISSUE=Neuroblastoma, and T-cell;
 RA Li W.B., Gruber C., Jessee J., Polayes D.;
 RT "Full-length cDNA libraries and normalization."
 RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A. (ISOFORM 3).
 RC TISSUE=Testis;
 RA Ottenwaelder B., Obermaier B., Mewes H.-W., Gassenhuber J.,
 RA Wiemann S.;
 RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A. (ISOFORM 1).
 RC TISSUE=Brain, and Lung;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Srausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaby S
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettman M., Madan A.C., Rodriguez S., Sanchez
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallus D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 human and mouse cDNA sequences."
 Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RL -!- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=3;
 CC Name=1;
 CC IsoId=Q86T03-1; Sequence=Displayed;
 CC Note=No experimental confirmation available;
 CC Name=2;
 CC IsoId=Q86T03-2; Sequence=VSP_007815;
 CC Note=No experimental confirmation available;
 CC

86T03-3; Sequence-VSP 007816, VSP_007817;
 y be due to intron retention.;

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 ail to license@isb-sib.ch).

 90; CAD61939.1; -;
 95; CAD62347.1; ALT INIT.
 97; CAB70896.1; ALT_INIT.
 97; AAH02867.2; -;
 97; AAH20947.1; -;
 T46382
 9299; C14orf9.
 protein; Transmembrane; Alternative splicing.
 POTENTIAL.
 112 234
 144 266
 47 47
 A -> AGKAPPO (in isoform 2).
 /FTID-VSP 007815.
 48 224
 APPFEGHFAVLPGEDPPYSLTSPDGSAPMITCRVCQ
 SLINVEGKHQHVKGVGCNEATPIKNAPPKKYVRCPCNC
 LLIKVTQRIACPRPKRIINLGVHPEPLSPFQPMGV
 RVICGHCKNTLWTEFTDLARCPCRKVSSIGRRYPRKR
 CICCFLGLLAV -> GKHPQGGKRGVAGPAGTLKAG
 EGAGPAAEAGPSRQVDCCTCDWRLPSLRNDRHSLGTGG
 SQPDRSANYEKPSELQGVQKQDPPTTVEHQWCK (in
 isoform 3).
 /FTID-VSP 007816.
 Missing (in isoform 3).
 /FTID-VSP 007817.
 P -> T (IN REF. 3; AAH20947).
 17 AA; 29469 MW; A85FE1F736366CBC CRC64;
 25 277
 208 208
 25; Score 7; DB 1; Length 277;
 arity 100.0%; Pred. No. 66;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 LAV 76
 |||||
 LAV 224
 STANDARD; PRT; 310 AA.
 1;
 (Rel. 41, Created)
 (Rel. 41, Last sequence update)
 (Rel. 43, Last annotation update)
 beta (LT-beta) (Tumor necrosis factor C) (TNF-C) (Tumor
 or ligand superfamily member 3).
 OR TNFC.
 (Woodchuck).
 atazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 heria; Rodentia; Sciurognathi; Scuridae; Sciurinae;
 995;
 4 N.A.
 4748; PubMed:10721723;
 all E.A., Brown C.L., Cullen J.M.;
 amphotoxin-alpha, -beta and tumor necrosis factor genes:
 characterization and biological activity.";
 -305(2000).
 : Cytokine that binds to LTBR/TNFRSF3. May play a specific
 immune response regulation. Provides the membrane anchor
 attachment of the heterotrimeric complex to the cell

CC -!- SUBUNIT: Heterotrimer of either two LTB and one LTA subunits
 CC (less prevalent) two LTA and one LTB subunits (By similarity)
 CC -!- SUBCELLULAR LOCATION: Type II membrane protein (Potential).
 CC -!- SIMILARITY: Belongs to the tumor necrosis factor family.
 CC
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 CC between the Swiss Institute of Bioinformatics and the EMBL out-
 CC the European Bioinformatics Institute. There are no restriction
 CC use by non-profit institutions as long as its content is in
 CC modified and this statement is not removed. Usage by and for
 CC entities requires a license agreement (See <http://www.isb-sib.ch>,
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; AF096268; AAF34866.1; -;
 CC EMBL; AF095587; AAF34865.1; -;
 CC HSSP; P01374; 1TNR.
 CC InterPro; IPR006053; TNF abc.
 CC InterPro; IPR006052; TNF family.
 CC InterPro; IPR008983; TNF_like.
 CC InterPro; IPR003636; TNF_subf.
 CC Pfam; PF00229; TNF; 1.
 CC PRINTS; PR01234; TNECROSISFCT.
 CC ProDom; PD002012; TNF_subf; 1.
 CC SMART; SM00207; TNF; 1.
 CC PROSITE; PS00251; TNF_1; FALSE_NEG.
 CC PROSITE; PS50049; TNF_2; 1.
 CC Cytokine; Transmembrane; Glycoprotein; Signal-anchor.
 FT DOMAIN 1 27
 FT SIGNAL-ANCHOR (TYPE-II MEMBRANE PRO-
 FT TRANSMEM 28 48
 FT (POTENTIAL).
 FT DOMAIN 49 310
 FT EXTRACELLULAR (POTENTIAL).
 FT CARBOHYD 272 272
 FT N-LINKED (GLCNAC...); (POTENTIAL).
 FT CONFLICT 280 280
 FT D -> H (IN REF. 1; AAF34865).
 SQ SEQUENCE 310 AA; 32644 MW; 73B354EFC8B3B3BE CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 310;
 Best Local Similarity 100.0%; Pred. No. 73;
 Matches 7; Conservative 0; Mismatches 0; Indels 0;
 QY 194 GLYLYYC 200
 |||||
 DB 181 GLYLYYC 187
 RESULT 49
 ISPH XANCP
 ID ISPH XANCP STANDARD; PRT; 316 AA.
 AC Q8P8G4;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE 4-hydroxy-3-methylbut-2-enyl diphosphate reductase (EC 1.17.1.2)
 GN ISPH OR LYTB OR KCC1157.
 OS Xanthomonas campestris (pv. campestris).
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
 OC Xanthomonadaceae; Xanthomonas.
 OC NCBI_TaxID=340;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 33913 / NCPPB 528;
 RX MEDLINE=22022145; PubMed=12024217;
 RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.J.
 RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N
 RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
 RA Camarotte G., Cannavan F., Cardozo J., Chambergo F., Ciapina L.P
 RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorriy H
 RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
 RA Fornighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
 RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.I.
 RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.
 RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
 RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V
 RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,

Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D., Antos M., Truffi D., Tsai S.M., White F.P., Kitajima J.P.;
the genomes of two Xanthomonas pathogens with differing ties";
-463(2002).
Converts 1-hydroxy-2-methyl-2-(E)-butenyl 4-diphosphate ntenyl diphosphate (IPP) and dimethylallyl diphosphate y similarity).
ACTIVITY: Isopentenyl diphosphate + NAD(P)(+) + H(2)O = oxy-3-methylbut-2-en-1-yl diphosphate + NAD(P)H.
onmevalonate terpenoid biosynthesis pathway; seventh P.
The genome sequence of the plant pathogen Xylella fastidiosa.";
Nature 406:151-159(2000).
[2]
SEQUENCE FROM N.A.
RP STRAIN=Temeculal / ATCC 700964;
RX MEDLINE=22421331; PubMed=12533478;
RA Van Sluys M.A., de Oliveira M.C., Monteiro-Vitorello C.B., Miyaki C.Y., Furlan L.R., Camargo L.E.A., da Silva A.C.R., Moon D. Takita M.A., Lemos E.G.M., Machado M.A., Ferro M.I.T., da Silva F. Goldman M.H.S., Goldman G.H., Lemos M.V.F., El-Dorri H., Tsai S.M. Carrer H., Carraro D.M., de Oliveira R.C., Nunes L.R., Siqueira W. Coutinho L.L., Kimura E.T., Ferro E.S., Harakava R., Kuramae E.E., Marino C.L., Gigliotti E., Abreu I.L., Alves L.M.C., do Amaral A.M. Baia G.S., Blanco S.R., Brito M.S., Cammavan F.S., Celestino A.V., da Cunha A.F., Fenille R.C., Ferro J.A., Formighieri E.F., Kishi I. Leoni S.G., Oliveira A.R., Rosa V.E. Jr., Sassaki F.T., Sena J.A.I. de Souza A.A., Truffi D., Tsukumo F., Yanai G.M., Zaros L.G., Civerolo E.L., Simpson A.J.G., Almeida N.F. Jr., Setubal J.C., Kitajima J.P.;
"Comparative analyses of the complete genome sequences of Pierce's disease and citrus variegated chlorosis strains of Xylella fastidiosa.";
J. Bacteriol. 185:1018-1026(2003).
CC -!- FUNCTION: Converts 1-hydroxy-2-methyl-2-(E)-butenyl 4-diphosphate into isopentenyl diphosphate (IPP) and dimethylallyl diphosphate (DMAPP) (By similarity).
CC -!- CATALYTIC ACTIVITY: Isopentenyl diphosphate + NAD(P)(+) + H(2) (E)-4-hydroxy-3-methylbut-2-en-1-yl diphosphate + NAD(P)H.
CC -!- PATHWAY: Nonmevalonate terpenoid biosynthesis pathway; seventh (last) step.
CC -!- SIMILARITY: Belongs to the ispH family.

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CC EMBL; AE004050; AAF85215.1; -;
DR EMBL; AE012558; AAO29279.1; -;
DR PIR; C82561; C82561.
DR HAMAP; MF_00191; -; 1.
DR InterPro; IPR003451; LysB.
DR Pfam; PF02401; LysB; 1.
DR TIGRFAMs; TIGR00216; isph lytB; 1.
KW Isoprene biosynthesis; Complete proteome; Oxidoreductase; NADP.
SQ SEQUENCE 316 AA; 34704 MW; 1A2E80B9A98D334A CRC64;

Query Match 2.5%; Score 7; DB 1; Length 316;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps
QY 217 LVDGVLA 223
DB 276 LVDGVLA 282
|||||
RESULT 51
ODPB_BACSU STANDARD; PRT; 324 AA.
ID ODPB_BACSU

RA Moon D.H., Nagai M.A., Nascimento A.L.T.O., Netto L.R.S., Nhani A. Jr., Nobrega F.G., Nunes L.R., Oliveira M.A., de Oliveira M.C., de Oliveira R.C., Palmieri D.A., Paris A., Peixoto B.R., Pereira G.A.G., Pereira H.A. Jr., Pesquero J.B., Raquiao R.B., Roberto P.G., Rodrigues V. de Rosa A.J.M., de Rosa V.E. Jr., de Sa R.G., Santelli R.V., Sawasaki H.E., da Silva A.C.R., da Silva A.M., da Silva F.R., Silva W.A. Jr., da Silveira J.F., Silvestri M.L.Z., Siqueira W.J., de Souza A.A., de Souza A.P., Terenzi M.F., Truffi D., Tsai S.M., Tshako M.H., Vallada H., Van Sluys M.A., Verjovski-Almeida S., Vettore A.L., Zago M.A., Zatz M., Meidanis J., Setubal J.C.;
"The genome sequence of the plant pathogen Xylella fastidiosa.";
Nature 406:151-159(2000).
[2]
SEQUENCE FROM N.A.
RP STRAIN=Temeculal / ATCC 700964;
RX MEDLINE=22421331; PubMed=12533478;
RA Van Sluys M.A., de Oliveira M.C., Monteiro-Vitorello C.B., Miyaki C.Y., Furlan L.R., Camargo L.E.A., da Silva A.C.R., Moon D. Takita M.A., Lemos E.G.M., Machado M.A., Ferro M.I.T., da Silva F. Goldman M.H.S., Goldman G.H., Lemos M.V.F., El-Dorri H., Tsai S.M. Carrer H., Carraro D.M., de Oliveira R.C., Nunes L.R., Siqueira W. Coutinho L.L., Kimura E.T., Ferro E.S., Harakava R., Kuramae E.E., Marino C.L., Gigliotti E., Abreu I.L., Alves L.M.C., do Amaral A.M. Baia G.S., Blanco S.R., Brito M.S., Cammavan F.S., Celestino A.V., da Cunha A.F., Fenille R.C., Ferro J.A., Formighieri E.F., Kishi I. Leoni S.G., Oliveira A.R., Rosa V.E. Jr., Sassaki F.T., Sena J.A.I. de Souza A.A., Truffi D., Tsukumo F., Yanai G.M., Zaros L.G., Civerolo E.L., Simpson A.J.G., Almeida N.F. Jr., Setubal J.C., Kitajima J.P.;
"Comparative analyses of the complete genome sequences of Pierce's disease and citrus variegated chlorosis strains of Xylella fastidiosa.";
J. Bacteriol. 185:1018-1026(2003).
CC -!- FUNCTION: Converts 1-hydroxy-2-methyl-2-(E)-butenyl 4-diphosphate into isopentenyl diphosphate (IPP) and dimethylallyl diphosphate (DMAPP) (By similarity).
CC -!- CATALYTIC ACTIVITY: Isopentenyl diphosphate + NAD(P)(+) + H(2) (E)-4-hydroxy-3-methylbut-2-en-1-yl diphosphate + NAD(P)H.
CC -!- PATHWAY: Nonmevalonate terpenoid biosynthesis pathway; seventh (last) step.
CC -!- SIMILARITY: Belongs to the ispH family.

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CC EMBL; AE004050; AAF85215.1; -;
DR EMBL; AE012558; AAO29279.1; -;
DR PIR; C82561; C82561.
DR HAMAP; MF_00191; -; 1.
DR InterPro; IPR003451; LysB.
DR Pfam; PF02401; LysB; 1.
DR TIGRFAMs; TIGR00216; isph lytB; 1.
KW Isoprene biosynthesis; Complete proteome; Oxidoreductase; NADP.
SQ SEQUENCE 316 AA; 34704 MW; 1A2E80B9A98D334A CRC64;

Query Match 2.5%; Score 7; DB 1; Length 316;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps
QY 217 LVDGVLA 223
DB 276 LVDGVLA 282
|||||
RESULT 51
ODPB_BACSU STANDARD; PRT; 324 AA.
ID ODPB_BACSU

Rel. 18, Created)
 Rel. 18, Last sequence update)
 Rel. 42, Last annotation update)
 drogenase E1 component, beta subunit (EC 1.2.4.1) (S
 Da subunit).
 OR BSU14590.
 ilis.
 micutes; Bacillales; Bacillaceae; Bacillus.
 23;
 N.A.
 558; PubMed=1697575;
 lva A., Paulin L., Arvidson S., Palva I.;
 complex of *Bacillus subtilis*: sequence analysis and
 yruvate dehydrogenase.";
 172:5052-5063(1990).
 N.A.
 187; PubMed=8969500;
 Aldwell R., Enfield L., Ferrari E.;
 E (124 degrees-127 degrees) region of the *Bacillus*
 chromosome: sequencing of a 27 kb segment and
 n of several genes in the area.";
 142:3033-3037(1996).
 N.A.
 Ferrari E.;
 lysis of the mobA-ampS region of the *Bacillus subtilis*
 TL-1997) to the EMBL/GenBank/DBJ databases.
 N.A.
 033; PubMed=9384377;
 sawara N., Moszer I., Albertini A.M., Alloni G.,
 artero M.G., Bessieres P., Bolotin A., Borchert S.,
 oursier L., Brans A., Braun M., Brignell S.C., Bron S.,
 Bruchet C.V., Caldwell B., Capuano V., Carter N.M.,
 dani J.J., Conerton I.F., Cummings N.J., Daniel R.A.,
 levine K.M., Dusterhoft A., Ehrlich S.D., Emerson P.T.,
 Errington J., Fabret C., Ferrari E., Foulger D.,
 ita M., Fujita Y., Fuma S., Galizzi A., Galleron N.,
 aser P., Goffeau A., Gollightly E.J., Grandi G.,
 Guy B.J., Haga K., Halech J., Harwood C.R., Henaut A.,
 Iolsappel S., Hosono S., Hullo M.F., Itaya M., Jones L.,
 amata D., Kasahara Y., Klaerr-Blanchard M., Klein C.,
 Koetter P., Koningsstein G., Krogh S., Kumano M.,
 pidus A., Lardinois S., Lauber J., Lazarevic V.,
 line A., Liu H., Masuda S., Maue C., Medigue C.,
 illado R.P., Mizuno M., Moestl D., Nakai S., Noback M.,
 Kelly M., Ogawa K., Ogiwara A., Oudega B., Park S.H.,
 il T.M., Portetelle D., Porwollik S., Prescott A.M.,
 Pujic P., Burnelle B., Rapoport G., Rey M., Reynolds S.,
 volta C., Rocha E., Roche B., Rose M., Sadaie Y.,
 ilan E., Schleich S., Schroeder R., Scoffone F.,
 Sekowska A., Seror S.J., Serro P., Shin B.S., Soldo B.,
 accioni E., Takagi T., Takahashi H., Takemaru K.,
 Tamakoshi A., Tanaka T., Terpstra P., Tognoni A.,
 chiya S., Vandebol M., Vannier F., Vassarotti A.,
 abutt R., Wedler E., Wedler H., Weitzenegger T.,
 Vipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,
 Yoshikawa H.F., Zumbstein E., Yoshikawa H., Danchin A.;
 a genome sequence of the Gram-positive bacterium *Bacillus*
 19-256(1997).
 : The pyruvate dehydrogenase complex catalyzes the overall
 n of pyruvate to acetyl-CoA and CO(2). It contains
 copies of three enzymatic components: pyruvate
 anase (E1), dihydrolipoamide acetyltransferase (E2) and

lipoamide dehydrogenase (E3).
 -!- FUNCTION: THE B.SUBTILIS PDH COMPLEX POSSESSES ALSO BRANCHED-
 2-OXOACID DEHYDROGENASE (BCDH) ACTIVITY.
 -!- CATALYTIC ACTIVITY: Pyruvate + lipoamide = S-
 acetyl-dihydrolipoamide + CO(2).
 -!- COFACTOR: Thiamine pyrophosphate.
 -!- SUBUNIT: Heterodimer of an alpha and a beta chain.
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 or send an email to license@isb-sib.ch).
 EMBL; M57435; AAA62682.1; --
 EMBL; AF012285; AAC24933.1; --
 EMBL; Z99111; CAB13332.1; --
 PIR; C36718; C36718.
 HSSP; P09061; 10S0.
 Subtilist; BG10208; pdhB.
 InterPro; IPR009014; Transketo_C_like.
 InterPro; IPR005476; Transketolase_C.
 InterPro; IPR005475; Transketolase_CR.
 Pfam; PF02779; transket_pyr; 1.
 Pfam; PF02780; transketolase_C; 1.
 Glycolysis; Oxidoreductase; Flavoprotein; Thiamine pyrophosphate;
 Complete proteome.
 INIT MET 0 BY SIMILARITY.
 SEQUENCE 324 AA; 35343 MW; D2A7C9B32DEDF0D CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 324;
 Best Local Similarity 100.0%; Pred. No. 75;
 Matches 7; Conservative 0; Mismatches 0; Indels 0;
 QY 59 LALGLGL 65
 DB 65 LALGLGL 71
 RESULT 52
 SRA6 CAEBL STANDARD; PRT; 329 AA.
 ID SRA6 CAEBL STANDARD; PRT; 329 AA.
 AC Q09208;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Serpentine receptor class alpha 6 (Sra-6 protein).
 GN SRA-6 OR AH6.10.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoi
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 OX NCBI_Taxid=62359;
 [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Bristol N2;
 RA Jassal B.;
 RL Submitted (JAN-1995) to the EMBL/GenBank/DBJ databases.
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
 CC -!- SIMILARITY: BELONGS TO THE C.ELEGANS RECEPTOR-LIKE PROTEIN SI
 FAMILY.
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 or send an email to license@isb-sib.ch).
 EMBL; Z48009; CAA88083.1; --
 PIR; T18619; T18619.

```

10; CS01451.
11; Sra; 1.
12; TMPTREINSRA.
13; Multigene family.
14; 46 POTENTIAL.
15; 124 POTENTIAL.
16; 163 POTENTIAL.
17; 207 POTENTIAL.
18; 258 POTENTIAL.
19; 293 POTENTIAL.
20; AA; 37951 MW; 7C0963ADA53A29F3 CRC64;
21; 2.5%; Score 7; DB 1; Length 329;
22; 100.0%; Pred. No. 76;
23; 0; Mismatches 0; Indels 0; Gaps 0;
24;
25; ARR 140
26; ||
27; ARR 303
28;
29; STANDARD; PRT; 335 AA.
30;
31; rel. 38, Created)
32; rel. 38, Last sequence update)
33; rel. 38, Last annotation update)
34; associated protein.
35;
36; .cus (Rat).
37; Azoo; Chordata; Craniata; Vertebrata; Euteleostomi;
38; Meria; Rodentia; Sciurognatha; Muridae; Rattus.
39; 16;
40; N.A.
41; TISSUE=Uterus;
42; A.; Lessing J.B., Kraicer P.F., Kidron T.;
43; associated uterine protein.;
44; 1999) to the EMBL/GenBank/DBJ databases.
45; R LOCATION: Integral membrane protein (Potential).
46; : Belongs to the OST3 family.
47;
48; Entry is copyright. It is produced through a collaboration
49; Swiss Institute of Bioinformatics and the EMBL outstation -
50; Bioinformatics Institute. There are no restrictions on its
51; profit institutions as long as its content is in no way
52; his statement is not removed. Usage by and for commercial
53; res a license agreement (See http://www.isb-sib.ch/announce/
54; il to license@isb-sib.ch).
55;
56; AAB63294.2; -
57; 106844; OST3 OST6.
58; 106663; Thioresox dom2.
59; OST3 OST6; 1.
60;
61; 6 26 POTENTIAL.
62; 15 205 POTENTIAL.
63; 0 230 POTENTIAL.
64; 71 291 POTENTIAL.
65; 11 321 POTENTIAL.
66; AA; 37992 MW; AFB9DB35F1A06091 CRC64;
67;
68; 2.5%; Score 7; DB 1; Length 335;
69; 100.0%; Pred. No. 78;
70; 0; Mismatches 0; Indels 0; Gaps 0;
71;
72; AV 76
73; ||
74; AV 198

```

```

RESULT 54
LEU3 THEAQ STANDARD; PRT; 344 AA.
AC P24038;
DT 01-MAR-1992 (Rel. 21, Created)
DT 01-MAR-1992 (Rel. 21, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE 3-isopropylmalate dehydrogenase (EC 1.1.1.85) (Beta-IPM dehydroge
DE (IMDH) (3-IPM-DH).
GN LEUB.
OS Thermus aquaticus.
OC Bacteria; Deinococcus-Thermus; Deinococci; Thermales; Thermaceae;
OC Thermus.
OX NCBI_TaxID=271;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=YTL;
RX MEDLINE=92041736; PubMed=1939005;
RA Kirino H., Oshima T.;
RT "Molecular cloning and nucleotide sequence of 3-isopropylmalate
RT dehydrogenase gene (leuB) from an extreme thermophile, Thermus
RT aquaticus YT-1".
RL J. Biochem. 109:852-857(1991).
CC -!- FUNCTION: Catalyzes the oxidation of 3-carboxy-2-hydroxy-4-
CC methylpentanoate (3-isopropylmalate) to 3-carboxy-4-methyl-2-
CC oxopentanoate. The product decarboxylates to 4-methyl-2-
CC oxopentanoate.
CC -!- CATALYTIC ACTIVITY: 3-carboxy-2-hydroxy-4-methylpentanoate +
CC NAD(+) = 3-carboxy-4-methyl-2-oxopentanoate + NADH.
CC -!- PATHWAY: Leucine biosynthesis; third step.
CC -!- SUBUNIT: Homodimer.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic.
CC -!- SIMILARITY: Belongs to the isocitrate and isopropylmalate
CC dehydrogenases family. LeuB subfamily 1.
CC
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CC between the Swiss Institute of Bioinformatics and the EMBL out
CC the European Bioinformatics Institute. There are no restriction
CC use by non-profit institutions as long as its content is in
CC modified and this statement is not removed. Usage by and for c
CC entities requires a license agreement (See http://www.isb-sib.ch/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; D10700; BAA01542.1; -
CC PIR; S41223; DETWIT.
CC HSSP; P00351; 1XAA.
CC HAMAP; MF_01033; 1
CC InterPro; IPR001804; IsoDH.
CC InterPro; IPR004429; LeuB.
CC Pfam; PF00180; isodh; 1.
CC TIGRFAMs; TIGR00169; leuB; 1.
CC PROSITE; PS00470; IDH IMDH; 1.
CC Oxidoreductase; Leucine biosynthesis; NAD.
CC SEQUENCE 344 AA; 36949 MW; 4F31A9444E826408 CRC64;
Query Match 2.5%; Score 7; DB 1; Length 344;
Best Local Similarity 100.0%; Pred. No. 80;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 248 SGLLAIR 254
DB 88 SGLLAIR 94
|||||
|||||
RESULT 55
LEU3 DEIRA STANDARD; PRT; 352 AA.
AC Q9RTH9;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE 3-isopropylmalate dehydrogenase (EC 1.1.1.85) (Beta-IPM dehydroge
DE (IMDH) (3-IPM-DH).

```

5. adiodurans.
 nococcus-Thermus; Deinococci; Deinococcales;
 e; Deinococcus.
 99;
 N.A.
 TCC 13939 / DSM 20539 / NCIB 9279;
 896; PubMed=10567266;
 en J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
 Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
 Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
 J., Lam P., McDonald L., Utterback T., Zalewski C.,
 Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
 Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
 nce of the radioresistant bacterium Deinococcus
 1"; 1577(1999).
 Catalyzes the oxidation of 3-carboxy-2-hydroxy-4-
 tanate (3-isopropylmalate) to 3-carboxy-4-methyl-2-
 oate. The product decarboxylates to 4-methyl-2-
 oate.
 ACTIVITY: 3-carboxy-2-hydroxy-4-methylpentanoate +
 3-carboxy-4-methyl-2-oxopentanoate + NADH.
 Leucine biosynthesis; third step.
 Homodimer (By similarity).
 AR LOCATION: Cytoplasmic (By similarity).
 Y: Belongs to the isocitrate and isopropylmalate
 nases family. LeuB subfamily 1.
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 ail to license@isb-sib.ch).
 9; AAF11333.1; -
 G75355.
 1XAA.
 33; -; 1.
 001804; Isodh.
 004429; Leub.
 ; isodh; 1.
 R00169; leub; 1.
 470; IDH-IMDH; 1.
 e; Leucine biosynthesis; NAD; Complete proteome.
 2 AA; 37598 MW; 8BAE0E347F2AFA29 CRC64;
 2.5%; Score 7; DB 1; Length 352;
 arity 100.0%; Pred.No. 81;
 onservative 0; Mismatches 0; Indels 0; Gaps 0;
 ALR 254
 |||||
 ALR 94
 STANDARD; PRT; 357 AA.
 Rel. 29, Created)
 Rel. 29, Last sequence update)
 Rel. 41, Last annotation update)
 sphatase (BC 3.1.3.9) (G6Pase) (G-6-Pase).
 (Mouse).
 tazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 heria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NCBI TaxID=10090;
 [1]
 RN SEQUENCE FROM N.A.
 RP TISSUE=Liver;
 RC MEDLINE=94012716; PubMed=8407995;
 RX Shelly L.L., Lei K.-J., Pan C.-J., Sakata S.F., Ruppert S.,
 RA Schutz G., Chou J.Y.;
 RT "Isolation of the gene for murine glucose-6-phosphatase, the enz
 RL deficient in glycogen storage disease type 1A.";
 RN J. Biol. Chem. 268:21482-21485(1993).
 [2]
 RN SEQUENCE OF 1-76 FROM N.A.
 RC STRAIN=129/SV; TISSUE=Liver;
 RX MEDLINE=97277298; PubMed=9115220;
 RA Streeter R.S., Svitek C.A., Chapman S., Greenbaum L.E., Taub R.,
 RA O'Brien R.M.;
 RT "A multicomponent insulin response sequence mediates a strong
 RT repression of mouse glucose-6-phosphatase gene transcription by
 RT insulin.";
 RL J. Biol. Chem. 272:11698-11701(1997).
 CC -!- FUNCTION: May be a single membrane channel protein acting bot
 CC a hydrolase and a translocase. It is the key enzyme in homeos
 CC regulation of blood glucose levels.
 CC -!- CATALYTIC ACTIVITY: D-glucose 6-phosphate + H(2)O = D-glucose
 CC phosphate.
 CC -!- PATHWAY: Gluconeogenesis and glycogenolysis; last step.
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Endoplasmic
 CC reticulum.
 CC -!- TISSUE SPECIFICITY: Liver and kidney.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a coll
 CC between the Swiss Institute of Bioinformatics and the EMBL out
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 CC modified and this statement is not removed. Usage by and for c
 CC entities requires a license agreement (See <http://www.isb-sib.ch/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; U00445; AAC52122.1; -
 CC EMBL; U91573; AAC53166.1; -
 CC PIR; A48589; A48589.
 CC MGI:95607; G6pc.
 CC InterPro; IPR008934; AcPase_VanPerase.
 CC Pfam; PF01569; PAP2; 1.
 CC SMART; SM00014; acidPPc; 1.
 CC Glycogen biosynthesis; Hydrolase; Transmembrane; Glycoprotein;
 CC Endoplasmic reticulum.
 CC TRANSMEM 30 46 POTENTIAL.
 CC TRANSMEM 59 75 POTENTIAL.
 CC TRANSMEM 153 169 POTENTIAL.
 CC TRANSMEM 211 227 POTENTIAL.
 CC TRANSMEM 296 312 POTENTIAL.
 CC TRANSMEM 333 349 POTENTIAL.
 CC CARBOHYD 96 96 N-LINKED (GLCNAC...) (BY SIMILARIT
 CC SITE 354 357 PREVENT SECRETION FROM ER (POTENTIAL
 CC SEQUENCE 357 AA; 40480 MW; 292F9FCE39582692 CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 357;
 Best Local Similarity 100.0%; Pred.No. 82;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G
 Qy 61 LGGLAL 67
 Db 269 LGGLAL 275
 |||||
 |||||
 RESULT 57
 G6PT RAT
 ID_G6PT RAT STANDARD; PRT; 357 AA.
 AC P43428;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)

el. 41, Last annotation update)
phatase (EC 3.1.3.9) (G6Pase) (G-6-Pase).
cus (Rat).
azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
16;

N.A.

19; PubMed=7860767;
in S., Chung E., Buikuisen W., Naji A., Taub R.A.;
if glucose-6-phosphatase gene and protein expression
tive response in proliferating liver and diabetes.";
t. 95:832-841(1995).

N.A.

-Dawley;
95; PubMed=8198588;
gaud D.M., El-Maghrabi M.R., Pan W., Subir M.,
a cDNA for the catalytic subunit of rat liver
phatase: regulation of gene expression in FAO hepatoma
in, dexamethasone and cAMP.";
ys. Res. Commun. 201:302-309(1994).

N.A.

-Dawley; TISSUE=Liver;
50; PubMed=8865366;
ajima H., Horikawa Y., Hamaguchi T., Yamasaki T.,
amba M., Hanafusa T., Matsuzawa Y.;
d distribution of glucose-6-phosphatase catalytic
ger RNA and its changes in the diabetic state.";
ol. Pathol. Pharmacol. 93:13-24(1996).
May be a single membrane channel protein acting both as
e and a translocase. It is the key enzyme in homeostatic
of blood glucose levels.
ACTIVITY: D-glucose 6-phosphate + H(2)O = D-glucose +

luconeogenesis and glycogenolysis; last step.

R LOCATION: Integral membrane protein. Endoplasmic

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AA474381.1; ALT_INIT.

AA419966.1; -.

BAA24348.1; -.

C2371.

08934; AcPase VanPerase.

00326; PA_PTPase.

PAP2; l.

; acidppc; l.

ntesis; Hydrolase; Transmembrane; Glycoprotein;

ticulum.

0 46 POTENTIAL.

9 75 POTENTIAL.

3 169 POTENTIAL.

1 227 POTENTIAL.

6 312 POTENTIAL.

3 349 POTENTIAL.

6 96 N-LINKED (GLCNAC. . .) (BY SIMILARITY).

4 357 PREVENT SECRETION FROM ER (POTENTIAL).

8 118 G -> V (IN REF. 2).

AA; 40555 MW; C44960E102F4244D CRC64;

2.5%; Score 7; DB 1; Length 357;

Best Local Similarity 100.0%; Pred. No. 82;
Matches 7; Conservative 0; Mismatches 0; Indels 0; G;

QY 61 LGLGLAL 67

Db 269 LGLGLAL 275

RESULT 58

PONI_RABIT

ID PONI_RABIT STANDARD; PRT; 358 AA.

AC P27170; Q9BGN1; Q9BGN2; Q9BGN3;

DT 01-AUG-1992 (Rel. 23, Created)

DT 01-AUG-1992 (Rel. 23, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Serum paraoxonase/arylesterase 1 (EC 3.1.1.2) (EC 3.1.1.8.1) (PONI 1);

DE (Serum arylalkylphosphatase 1) (A-esterase 1) (Aromatic esterase

GN PONI OR PON.

OS Oryctolagus cuniculus (Rabbit).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.

OX NCBI_TaxID=9986;

RN [1]

RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.

RC TISSUE=Liver;

RX MEDLINE=92031445; PubMed=1657140;

RA Hassett C., Richter R.J., Humbert R., Chapline C., Crabb J.W.,

RA Omiecinski C.J., Furlong C.E.;

RT "Characterization of cDNA clones encoding rabbit and human serum

RT paraoxonase: the mature protein retains its signal sequence.";

RL Biochemistry 30:10141-10149(1991).

RN [2]

RP SEQUENCE FROM N.A., AND CHARACTERIZATION.

RX MEDLINE=93345100; PubMed=8393745;

RA Furlong C.E., Costa L.G., Hassett C., Richter R.J.,

RA Sundstrom J.A., Adler D.A., Distche C.M., Omiecinski C.J.,

RA Chapline C., Crabb J.W.;

RT "Human and rabbit paraoxonases: purification, cloning, sequencing

RT mapping and role of polymorphism in organophosphate detoxification

RL Chem. Biol. Interact. 87:35-48(1993).

RN [3]

RP SEQUENCE FROM N.A., FUNCTION, AND VARIANTS SER-81; GLU-92 AND GLY.

RC STRAIN=New Zealand white; TISSUE=Liver;

RX MEDLINE=21163843; PubMed=11266077;

RA Watson C.E., Draganov D.I., Billecke S.S., Bisgaier C.L., La Du B

RT "Rabbits possess a serum paraoxonase polymorphism similar to the l

RT Q192R.";

RL Pharmacogenetics 11:123-134(2001).

RN [4]

RP CHARACTERIZATION, AND SEQUENCE OF 1-20.

RX MEDLINE=92031444; PubMed=1718413;

RA Furlong C.E., Richter R.J., Chapline C., Crabb J.W.;

RT "Purification of rabbit and human serum paraoxonase.";

RL Biochemistry 30:10133-10140(1991).

CC -!- FUNCTION: Hydrolyzes the toxic metabolites of a variety of

CC organophosphorus insecticides. Capable of hydrolyzing a broad

CC spectrum of organophosphate substrates and a number of aromati

CC carboxylic acid esters. Mediates an enzymatic protection of l

CC density lipoproteins against oxidative modification.

CC -!- CATALYTIC ACTIVITY: Aryl dialkyl phosphate + H(2)O = dialkyl

CC phosphate + an aryl alcohol.

CC -!- SUBCELLULAR LOCATION: Extracellular.

CC -!- TISSUE SPECIFICITY: Plasma.

CC -!- PTM: Glycosylated.

CC -!- PTM: The signal sequence is not cleaved.

CC -!- POLYMORPHISM: There are two allelic forms, allozyme A and B, v

CC differ in their substrate specificity. Both forms have simila;

CC arylesterase activity but allozyme B possesses greater paraox

CC activity. Allozyme A is better at protecting LDL from oxidati

CC -!- SIMILARITY: Belongs to the paraoxonase family.

CC -!- CAUTION: Ref.3 (AAK06398) sequence differs from that shown due

CC a stop codon in position 355.

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AAA31452.1; -;
AA227713.2; -;
1; AAK06398.1; ALT_TERM.
2; AAK06399.1; -;
3; AAK06400.1; -;
B40354.
6; C:extracellular; NAS.
9; P:antioxidant activity; IDA.
3; P:arylalkylphosphatase activity; NAS.
4; P:arylesterase activity; IDA.
3; P:response to organophosphorus; IDA.
002640; Arylesterase.
008363; Paraoxonase1.
008364; Paraoxonase2.
; Arylesterase; 1.
85; PARAOXONASE.
86; PARAOXONASE1.
87; PARAOXONASE2.
tioxidant; Glycoprotein; Signal; Multigene family;

0 0
1 ?
41 352
BY SIMILARITY.
49 49 N-LINKED (GLCNAC. .) (POTENTIAL).
52 252 N-LINKED (GLCNAC. .) (POTENTIAL).
69 269 N-LINKED (GLCNAC. .) (POTENTIAL).
23 323 N-LINKED (GLCNAC. .) (POTENTIAL).
81 81 P -> S (IN ALLELE A).
92 92 K -> E (IN ALLELE A).
100 100 S -> G (IN ALLELE A).
66 66 A -> V (IN REF. 3).
19 319 A -> V (IN REF. 3); AAK06398).
8 AA; 39878 MW; C40C45E5F6E5E5FDF CRC64;

2.5%; Score 7; DB 1; Length 358;
arity 100.0%; Pred.No. 82;
conservative 0; Mismatches 0; Indels 0; Gaps 0;

LAL 67
|||
LAL 15

STANDARD; PRT; 365 AA.

Rel. 35, Created)
Rel. 35, Last sequence update)
Rel. 43, Last annotation update)
oxidoreductase chain 8 (EC 1.6.99.5) (NADH dehydrogenase
(NDH-1, chain 8)).

ophilus.
nococcus-Thermus; Deinococci; Thermales; Thermaceae;
4;

1 N.A.
ATCC 27634;
490; PubMed=9020134;
S.S., Sled, V.D., Ohnishi T., Yagi T.;
translocating NADH-quinone oxidoreductase (NDH-1) of
bacterium Thermus thermophilus HB-8. Complete DNA

RT sequence of the gene cluster and thermostable properties of the
expressed NQO2 subunit.";
J. Biol. Chem. 272:4201-4211 (1997).
-!- FUNCTION: NDH-1 shuttles electrons from NADH, via FMN and iron
sulfur (Fe-S) centers, to quinones in the respiratory chain.
Immediate electron acceptor for the enzyme in this species is
believed to be menaquinone. Couples the redox reaction to proton
translocation (for every two electrons transferred, four protons
ions are translocated across the cytoplasmic membrane), and thus
conserves the redox energy in a proton gradient.
-!- CATALYTIC ACTIVITY: NADH + quinone = NAD(+) + quinol.
-!- SUBUNIT: COMPOSED OF 14 DIFFERENT SUBUNITS. SUBUNIT NQO7-14
CONSTITUTE THE MEMBRANE SECTOR OF THE COMPLEX.
-!- SUBCELLULAR LOCATION: Integral membrane protein.
-!- SIMILARITY: Belongs to the complex I subunit 1 family.

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modified and this statement is not removed. Usage by and for c
entities requires a license agreement (See <http://www.isb-sib.ch/>
or send an email to license@isb-sib.ch).

CC EMBL; U52917; AAA97945.1; -;
DR PIR; T11905; T11905.
DR InterPro; IPR001694; Resp_NADH_dh1.
DR Pfam; PF00146; NADHdh; 1.
DR PROSITE; PS00667; COMPLEX1_NDI_1; 1.
DR PROSITE; PS00668; COMPLEX1_NDI_2; 1.
KW Oxidoreductase; NAD; Quinone; Transmembrane.
FT TRANSMEM 11 31 POTENTIAL.
FT TRANSMEM 80 100 POTENTIAL.
FT TRANSMEM 120 140 POTENTIAL.
FT TRANSMEM 157 177 POTENTIAL.
FT TRANSMEM 192 212 POTENTIAL.
FT TRANSMEM 252 272 POTENTIAL.
FT TRANSMEM 273 293 POTENTIAL.
FT TRANSMEM 310 330 POTENTIAL.
FT TRANSMEM 336 356 POTENTIAL.
SQ SEQUENCE 365 AA; 41008 MW; AE920CC029333C09 CRC64;

Query Match 2.5%; Score 7; DB 1; Length 365;
Best Local Similarity 100.0%; Pred.No. 84;
Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 61 LGLGLAL 67
|||
Db 164 LGLGLAL 170

RESULT 60

BENE ACICA
ID BENE ACICA STANDARD; PRT; 394 AA.
AC P07775;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Benzoate membrane transport protein.
GN BENE.
OS Acinetobacter calcoaceticus.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Moraxellaceae; Acinetobacter.
OX NCBI_TaxID=471;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BD413 / ADP1;
RX MEDLINE=91358314; PubMed=1885518;
RA Neidle E.L., Hartnett C., Ornston N.L., Bairoch A., Rekik M.,
RA Harayama S.;
FT "Nucleotide sequences of the Acinetobacter calcoaceticus benABC
RT for benzoate 1,2-dioxygenase reveal evolutionary relationships an
multicomponent oxygenases.";

173-5385-5395(1991).
PROBABLY INVOLVED IN THE TRANSPORT OF BENZOATE.
R LOCATION: Integral membrane protein.
OUS: THE BEN OPERON ENCODE THE PROTEINS RESPONSIBLE FOR
ATION OF BENZOATE TO CATECHOL.

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il to license@isb-sib.ch).

; AAC46440.1; -
23481.
04711; BenE.
BenE; 1.
53; BenE; 1.
00843; BenE; 1.
matic hydrocarbons catabolism; Transmembrane.
AA: 42288 MW; 5466C4D460784BB5 CRC64;
2.5%; Score 7; DB 1; Length 394;
rity 100.0%; Pred. No. 90;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
LV 57
||
LV 88

STANDARD; PRT; 396 AA.
el. 38, Created)
el. 38, Last sequence update)
el. 42, Last annotation update)
g protein precursor 1 (DHH-1) (Cephalic hedgehog
[H].
(African clawed frog).
azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
achia; Anura; Mesobatrachia; Pipidae; Pipidae;
enopus.
5;
N.A.

52; PubMed=7671800;
Grew L.L., Lai C.-J., Lee J.J., von Kessler D.P.,
Chy P.A.;
ession and shared activities of members of the hedgehog
Xenopus laevis".
1:2337-2347(1995).
AL ECTODERM, NERVOUS SYSTEM AND SOMITES. INDUCES ECTOPIC
ND FORMATION IN EMBRYOS
R LOCATION: THE C-TERMINAL PEPTIDE DIFFUSES FROM THE
E THE N-TERMINAL PEPTIDE REMAINS ASSOCIATED WITH THE
CE. HEDGEHOG PROTEIN IS ALSO SECRETED IN EITHER CLEAVED
ED FORM TO MEDIATE SIGNALING TO OTHER CELLS (BY
).
TAL STAGE: DETECTABLE WITHIN THE EARLY GASTRULA. IN THE
PRESSION BECOMES RESTRICTED TO ANTERIOR STRUCTURES,
NG BOTH NEURAL PLATE AND ENDODERMAL CELLS.
-terminal domain displays an autophosphorylation activity
esterol transferase activity. Both activities result in
ge of the full-length protein and covalent attachment of
rol moiety to the C-terminal of the newly generated N-
ragment (N-product). This covalent modification appears
essential role in restricting the spatial distribution

of the protein activity to the cell surface. The N-product is
active species in both local and long-range signaling, whereas:
C-product has no signaling activity (By similarity).
-!- SIMILARITY: Belongs to the hedgehog family.

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modified and this statement is not removed. Usage by and for c
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or send an email to license@isb-sib.ch).

EMBL: U26349; AAA85163.1; -
DR HSP; O62226; 1VHH.
DR MEROPS; C46 UPW; -
DR InterPro; IPR009045; Hedgehog/DD_pept.
DR InterPro; IPR003587; Hedgehog hint_N.
DR InterPro; IPR003586; Hedgehog hint_C.
DR InterPro; IPR000320; HH signal.
DR InterPro; IPR006141; Intein S.
DR InterPro; IPR001767; Pept C46 hint.
DR InterPro; IPR001657; Peptidase_C46.
DR Pfam; PF01085; HH signal; 1.
DR Pfam; PF01079; Hint; 1.
DR PRINTS; PR00632; SONICHHOG.
DR ProDom; PD003042; HH signal; 1.
DR SMART; SM00305; HintC; 1.
DR SMART; SM00306; HintN; 1.
DR PROSITE; PS00817; INTEIN_N_TER; 1.
KW Developmental protein; Autocatalytic cleavage; Hydrolase; Proteas
Signal; Lipoprotein; Palmitate.
FT SIGNAL 1 22 POTENTIAL.
FT CHAIN 23 396 DESERT HEDGEHOG PROTEIN 1.
FT CHAIN 23 197 DESERT HEDGEHOG PROTEIN 1 N-PRODUCT.
FT CHAIN 198 396 DESERT HEDGEHOG PROTEIN 1 C-PRODUCT.
FT DOMAIN 276 279 POLY-SER.
FT SITE 197 198 CLEAVAGE (AUTO-) (BY SIMILARITY).
FT SITE 267 267 INVOLVED IN AUTO-CLEAVAGE (BY
SIMILARITY).
FT ACT_SITE 270 270 ESSENTIAL FOR AUTO-CLEAVAGE (BY
SIMILARITY).
FT LIPID 23 23 N-palmitoyl cysteine (By similarity)
FT LIPID 197 197 Cholesterol glycine ester (By
similarity).
FT SEQUENCE 396 AA; 44087 MW; 774A3EC2268A5EE9 CRC64;
Query Match 2.5%; Score 7; DB 1; Length 396;
Best Local Similarity 100.0%; Pred. No. 90;
Matches 7; Conservative 0; Mismatches 0; Indels 0; G
QY 216 LLVDGVL 222
Db 331 LLVDGVL 337
|||||
RESULT 62
DHH2 XENLA
ID DHH2 XENLA STANDARD; PRT; 398 AA.
AC Q91611;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Desert hedgehog protein precursor 2 (DHH-2) (Hedgehog protein 4) (
DE HH4).
GN HH4.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.

852; PubMed=7671800;
McGrew L.L., Lai C.-J., Lee J.J., von Kessler D.P.,
Bachy P.A.;
expression and shared activities of members of the hedgehog
of *Xenopus laevis*.;
21:2337-2347(1995).
SIGNAL INVOLVED IN THE EARLY INDUCTION AND PATTERNING OF
SAL ECTODERM, NERVOUS SYSTEM AND SOMITES. INDUCES ECTOPIC
AND FORMATION IN EMBRYOS.
AR LOCATION: THE C-TERMINAL PEPTIDE DIFFUSES FROM THE
LE THE N-TERMINAL PEPTIDE REMAINS ASSOCIATED WITH THE
FACE. HEDGEHOG PROTEIN IS ALSO SECRETED IN EITHER CLEAVED
VED FORM TO MEDIANE SIGNALING TO OTHER CELLS (BY
Y).
C-terminal domain displays an autophosphorylation activity
lesterol transferase activity. Both activities result in
age of the full-length protein and covalent attachment of
erol moiety to the C-terminal of the newly generated N-
fragment (N-product). This covalent modification appears
in essential role in restricting the spatial distribution
rotein activity to the cell surface. The N-product is the
pecies in both local and long-range signaling, whereas the
has no signaling activity (By similarity).
Y: Belongs to the hedgehog family.

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AA085164.1; -
1VHH.
IPW; -
009045; Hedgehog/DD_pept.
003587; Hedgehog hint N.
003586; Hedgehog hintC.
000320; HH signal.
001767; Pept C46 hint.
001657; Peptidase_C46.
; HH signal; 1.
; Hint; 1.
32; SONICHHOG.
042; HH signal; 1.
5; HintG; 1.
6; HintN; 1.
- protein; Autocatalytic cleavage; Hydrolase; Protease;
- protein; Palmitate.
1 23 POTENTIAL.
24 398 DESERT HEDGEHOG PROTEIN 2.
24 199 DESERT HEDGEHOG PROTEIN 2 N-PRODUCT.
200 398 DESERT HEDGEHOG PROTEIN 2 C-PRODUCT.
278 281 POLY-SER.
199 200 CLEAVAGE (AUTO-) (BY SIMILARITY).
269 269 INVOLVED IN AUTO-CLEAVAGE (BY
SIMILARITY).
272 272 ESSENTIAL FOR AUTO-CLEAVAGE (BY
SIMILARITY).
24 24 N-palmitoyl cysteine (By similarity).
199 199 Cholesterol glycine ester (By
similarity).
98 AA; 44458 MW; DBC23AF85F69DD08 CRC64;
2.5%; Score 7; DB 1; Length 398;
larity 100.0%; Pred.No.90;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CGVL 222
|||
CGVL 339

RESULT 63
SELP_BOVIN
ID SELP_BOVIN STANDARD; PRT; 402 AA.
AC P49907; O19003;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Selenoprotein P-like protein precursor.
DE Selenoprotein P-like protein precursor.
GN SEPL1 OR SELP.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OC NCBI_TaxID=9913;
RN [1]
SEQUENCE FROM N.A.
RP TISSUE=Cerebellum;
RC MEDLINE=95364621; PubMed=7637580;
RA Saijoh K., Saito N., Lee M.J., Fujii M., Kobayashi T., Sumino K.
RT "Molecular cloning of cDNA encoding a bovine selenoprotein P-like
protein containing 12 selenocysteines and a (His-Pro) rich domain
insertion, and its regional expression.";
RT Brain Res. Mol. Brain Res. 30:301-311(1995).
RL [2]
SEQUENCE FROM N.A.
RP MEDLINE=98019090; PubMed=9358058;
RA Fujii M., Saijoh K., Kobayashi T., Fujii S., Lee M.J., Sumino K.
RT "Analysis of bovine selenoprotein P-like protein gene and availa
of metal responsive element (MRE) located in its promoter.";
RL Gene 199:211-217(1997).
CC -!- FUNCTION: It constitutes a major selenium pool in the brain;
may play an important role in developing and/or modulating t
morphology of neurons and/or glial cells.
CC -!- SUBCELLULAR LOCATION: Secreted (By similarity).
CC -!- TISSUE SPECIFICITY: Brain and kidney. Most prominently expre
in the cerebellar cortex, hippocampus and olfactory bulb.
CC -!- MISCELLANEOUS: The selenocysteines are all encoded by the op
codon, UGA.

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EMBL; D25220; BAA04949.2; -
EMBL; D88033; BAA23414.1; -
EMBL; D88031; BAA23414.1; JOINED.
EMBL; D88032; BAA23414.1; JOINED.
InterPro; IPR007672; Selp_C.
InterPro; IPR007671; Selp_N.
Pfam; PF04593; Selp_C; 1.
Pfam; PF04592; Selp_N; 1.
KW Glycoprotein; Signal; Selenium; Selenocysteine; Repeat.
SIGNAL 1 19
BY SIMILARITY.
SELENOPROTEIN P-LIKE PROTEIN.
FT CHAIN 20 402
FT SE_CYS 59 59
FT SE_CYS 297 297
FT SE_CYS 307 307
FT SE_CYS 338 338
FT SE_CYS 350 350
FT SE_CYS 363 363
FT SE_CYS 365 365
FT SE_CYS 372 372
FT SE_CYS 388 388
FT SE_CYS 390 390
FT SE_CYS 397 397
FT SE_CYS 399 399
FT SE_CYS 204 239
FT DOMAIN
H-P REPEATS.

0 266 POLY-HIS.
 8 181 SRPQ -> KALE (IN REF. 2).
 6 256 T -> P (IN REF. 2).
 2 282 L -> V (IN REF. 2).
 2 312 Y -> D (IN REF. 2).
 AA: 45018 MW; B7CF18751F808EFF CRC64;
 2.5%; Score 7; DB 1; Length 402;
 100.0%; Pred. No. 91;
 unservative 0; Mismatches 0; Indels 0; Gaps 0;
 LA 68
 LA 10
 STANDARD; PRT; 412 AA.
 el. 34, Created)
 el. 34, Last sequence update)
 el. 41, Last annotation update)
 te kinase, plasmid (EC 2.7.2.3).
 trophus (Ralstonia eutropha).
 asmid pHG1.
 eobacteria; Betaproteobacteria; Burkholderiales;
 ae; Ralstonia.
 N.A.
 DSM 428 / ATCC 17699;
 15; PubMed=7763137;
 J., Yoo J.-G., Bowien B.;
 the genes forming the distal parts of the two cbb CO2
 us from Alcaligenes eutrophus.";
 1. 163:291-299(1995).
 ACTIVITY: ATP + 3-phospho-D-glycerate = ADP + 3-
 glyceroyl phosphate.
 Calvin cycle.
 monomer (By similarity).
 R LOCATION: Cytoplasmic.
 : Belongs to the phosphoglycerate kinase family.
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 res a license agreement (see <http://www.isb-sib.ch/announce/>
 ul to license@isb-sib.ch).
 AAC43447.1; -.
 1PHP.
 15; -. 1.
 001576; PGK.
 PGK; 1.
 7; PGLYCKINASE.
 11; PGLYCERATE_KINASE; 1.
 Kinase; Calvin cycle; Plasmid.
 2 AA; 42298 MW; 9584C666859F7274 CRC64;
 2.5%; Score 7; DB 1; Length 412;
 100.0%; Pred. No. 93;
 unservative 0; Mismatches 0; Indels 0; Gaps 0;
 TAA 235
 TAA 293

PGKC_ALCEU
 ID PGKC_ALCEU STANDARD; PRT; 413 AA.
 AC P50319;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Phosphoglycerate kinase, chromosomal (EC 2.7.2.3).
 GN CBBKC.
 OS Alcaligenes eutrophus (Ralstonia eutropha).
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
 OC Burkholderiaceae; Ralstonia.
 OX NCBI_TaxID=510;
 RN [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=H16 / DSM 428 / ATCC 17699;
 RX MEDLINE=95283415; PubMed=7763137;
 RA Schaeferfohann J., Yoo J.-G., Bowien B.;
 RT "Analysis of the genes forming the distal parts of the two cbb CO:
 fixation operons from Alcaligenes eutrophus.";
 RL Arch. Microbiol. 163:291-299(1995).
 CC -|- CATALYTIC ACTIVITY: ATP + 3-phospho-D-glycerate = ADP + 3-
 phospho-D-glyceroyl phosphate.
 CC -|- PATHWAY: Calvin cycle.
 CC -|- SUBUNIT: Monomer (By similarity).
 CC -|- SUBCELLULAR LOCATION: Cytoplasmic.
 CC -|- SIMILARITY: Belongs to the phosphoglycerate kinase family.
 CC
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 or send an email to license@isb-sib.ch).
 EMBL; U12422; AAC43444.1; -.
 DR PIR; I39551; I39551.
 DR HSSP; P18912; 1PHP.
 DR HAMAP; MF_00145; -. 1.
 DR InterPro; IPR001576; PGK.
 DR Pfam; PF00162; PGK; 1.
 DR PRINTS; PR00477; PGLYCKINASE
 DR PROSITE; PS00111; PGLYCERATE_KINASE; 1.
 KW Transferase; Kinase; Calvin cycle.
 SQ SEQUENCE 413 AA; 42283 MW; 6B4C9D195566A90D CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 413;
 Best Local Similarity 100.0%; Pred. No. 93;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G:
 QY 229 EFSATAA 235
 |||||
 DB 288 EFSATAA 294
 RESULT 66
 NH59 CAEEL STANDARD; PRT; 416 AA.
 ID NH59 CAEEL
 AC QSTXJ1; Q9GTF2;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Nuclear hormone receptor family member nhr-59.
 GN NHR-59 OR T27B7.1.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoid;
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 OX NCBI_TaxID=6239;
 RN [1]
 RN SEQUENCE FROM N.A.
 RA Bogan A., Maina C.V., Yamamoto K., Cohen P., Sluder A.E.;
 RT "Caenorhabditis elegans nuclear receptor sequences exhibit biophy
 compatibility with the ligand-binding domain fold.";

Y-2000) to the EMBL/GenBank/DBJ databases.
 N.A.
 1 N2;
 sley P., O'Brien D.;
 G-1997) to the EMBL/GenBank/DBJ databases.
 N-2002) to the EMBL/GenBank/DBJ databases.
 Orphan nuclear receptor.
 AR LOCATION: Nuclear (Potential).
 Y: Belongs to the nuclear hormone receptor family.
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 ail to license@isb-sib.ch).
 7; AAG15156.1; -;
 5; AAF02172.2; -;
 1AGY.
 7.1; CE24017.
 008946; Str ncl receptor.
 001628; Znf_C4steroid.
 ; Zf-C4; 1.
 47; STROLDINGER.
 035; Znf_C4steroid; 1.
 9; Znf_C4; 1.
 031; NUCLEAR RECEPTOR; FALSE NEG.
 nscription regulation; DNA-Binding; Nuclear protein;
 20 87 NUCLEAR RECEPTOR-TYPE.
 20 40 C4-TYPE.
 57 82 C4-TYPE.
 6 AA; 47721 MW; D4A21CA587ED96D3 CRC64;
 2.5%; Score 7; DB 1; Length 416;
 arity 100.0%; Pred.No. 94;
 onservative 0; Mismatches 0; Indels 0; Gaps 0;
 ISA 10
 ||||
 ISA 177
 STANDARD; PRT; 418 AA.
 2;
 Rel. 09, Created)
 Rel. 35, Last sequence update)
 Rel. 35, Last annotation update)
 ke protease inhibitor 6 precursor (CPI-26) (Serine
 bitor 3) (SPI-3) (SPI-2.2).
 ricus (Rat)
 itazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 heria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 116;
 1 N.A.
 1305; PubMed=1864837;
 ata S., Misumi Y., Takami N., Ikehara Y.;
 .oning and characterization of rat contrapsin-like
 .bitor and related proteins.;
 .09:243-250(1991).
 .1-408 FROM N.A.

RX MEDLINE=90306038; PubMed=1694763;
 RA Pages G., Rouayrenc J.F., le Cam G., Mariller M., le Cam A.;
 RT "Molecular characterization of three rat liver serine-protease
 RT inhibitors affected by inflammation and hypophysectomy. Protein
 RT mRNA analysis and cDNA cloning.";
 RL Eur. J. Biochem. 190:385-391(1990).
 RN [3]
 RP SEQUENCE OF 203-408 FROM N.A.
 RX MEDLINE=87144617; PubMed=3493437;
 RA Hill R.E., Hastie N.D.;
 RT "Accelerated evolution in the reactive centre regions of serine
 RT protease inhibitors.";
 RL Nature 326:96-99(1987).
 CC -!- FUNCTION: INHIBITS TRYPSIN, BUT NOT CHYMOTRYPSIN OR
 CC ELASTASE.
 CC -!- TISSUE SPECIFICITY: Liver.
 CC -!- INDUCTION: By acute inflammation.
 CC -!- SIMILARITY: Belongs to the serpin family.
 CC -!- CAUTION: It is uncertain whether Met-1 or Met-11 is the initi
 CC This SWISS-PROT entry is copyright. It is produced through a coll
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 CC entities requires a license agreement (See <http://www.isb-sib.ch/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; D00753; BRA00650.1; -;
 CC EMBL; X16359; CAA34408.1; -;
 CC EMBL; X13150; CAA31548.1; -;
 CC HSSP; P01011; 2ACH.
 CC InterPro; IPR000215; Serpin.
 CC SMART; SM00093; SERPIN; 1.
 CC PROSITE; PS00284; SERPIN; 1.
 KW Serpin; Serine protease inhibitor; Glycoprotein; Signal.
 FT SIGNAL 1 29
 FT CHAIN 30 418
 FT ACT SITE 381 382
 FT REACTIVE BOND (POTENTIAL).
 FT CONTRAPSIN-LIKE PROTEASE INHIBITOR {
 FT N-LINKED (GLCNAc. . .) (POTENTIAL).
 FT N-LINKED (GLCNAc. . .) (POTENTIAL).
 FT N-LINKED (GLCNAc. . .) (POTENTIAL).
 FT A -> D (IN REF. 2).
 FT C -> S (IN REF. 2).
 FT D -> H (IN REF. 2).
 FT N -> S (IN REF. 2).
 FT P -> S (IN REF. 2).
 FT E -> EE (IN REF. 3).
 FT V -> A (IN REF. 3).
 FT S -> F (IN REF. 2).
 FT Q -> E (IN REF. 3).
 FT M -> V (IN REF. 3).
 SQ SEQUENCE 418 AA; 46651 MW; AADEF087190B44F CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 418;
 Best Local Similarity 100.0%; Pred.No. 94;
 Matches 7; Conservative 0; Mismatches 0; Indels 0;
 QY 74 LAVVSLG 80
 Db 82 LAVVSLG 88
 RESULT 68
 ENO_PYRAE STANDARD; PRT; 419 AA.
 ID ENO_PYRAE
 AC Q82YE7;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Enolase (EC 4.2.1.11) (2-phosphoglycerate dehydratase) (2-phosph
 DE glycerate hydro-lyase).

SS71103.
 41916; --
 ; YUR084W.
 0; C:signalosome complex; IDA.
 4; P:adaptation to pheromone during conjugation . . .; IMP.
 00017; PCI.
 ; PCI; 1.
 protein.
 3 AA; 49482 MW; 750CDA631916A621 CRC64;
 2.5%; Score 7; DB 1; Length 423;
 arity 100.0%; Pred. No. 95;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;
 LAL 61
 ||||
 LAL 272
 STANDARD; PRT; 424 AA.
 Rel. 11, Created
 Rel. 35, Last sequence update)
 Rel. 39, Last annotation update)
 a sperm-binding protein 3 precursor (Zona pellucida
 ZP3) (sperm receptor) (Zona pellucida protein C).
 (Mouse).
 -tazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 heria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 090;
 (N.A.
 926; PubMed=3378665;
 ; Chamberlin M.E.; Baur A.W.; Sobieski D.A.; Dean J.;
 analysis of cDNA coding for ZP3, a sperm binding protein
 zona pellucida.";
 7:287-295 (1988).
 87.
 V-1996) to the EMBL/GenBank/DBJ databases.
 (N.A.
 TISSUE=Liver;
 048; PubMed=2541416;
 Wassarman P.M.;
 sequence of the gene encoding zona pellucida glycoprotein
 use sperm receptor.";
 Res. 17:2861-2863 (1989).
 (N.A.
 451; PubMed=2842770;
 Roller R.J.; Fimiani C.M.; Wassarman D.A.;
 ;
 cture of the mouse sperm receptor polypeptide determined
 oning.";
 acad. Sci. U.S.A. 85:6409-6413 (1988).
 (9-63; 197-204; 219-233 AND 261-275.
 9795; PubMed=1330788;
 Wassarman P.M.;
 on of a region of mouse zona pellucida glycoprotein mzp3
 is sperm receptor activity.";
 14:309-317 (1992).
 ; Functions as a sperm-receptor. It is responsible for
 sion to the zona pellucida, and may contribute to the
 specificity of the insemination.
 ZP3 FORMS WITH ZP1 AND ZP2 THE ZONA PELLUCIDA, IN
 ; AND ZP3 COMPLEX INTO COPOLYMERS CROSS-LINKED BY ZP1.

CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Extracellular
 CC matrix.
 CC -!- TISSUE SPECIFICITY: Oocytes.
 CC -!- DEVELOPMENTAL STAGE: Expressed during the 2-week growth phase
 CC oogenesis, prior to ovulation.
 CC -!- PM: Sulfated glycoprotein with O-linked oligosaccharides.
 CC -!- SIMILARITY: Contains 1 ZP domain.
 CC -----
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 CC between the Swiss Institute of Bioinformatics and the EMBL out
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 CC entities requires a license agreement (See <http://www.isb-sib.ch/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M20026; AAB18629.1; --
 CC EMBL; X14376; CAA32550.1; --
 CC PIR; A30334; A30334.
 CC MGD; MGI:99215; Zp3.
 CC InterPro; IPR001507; Endoglin/CD105.
 CC Pfam; PF00100; zona_pellucida; 1.
 CC PRINTS; PR00023; ZPELLUCIDA.
 CC SMART; SM00241; ZP; 1; DOMAIN; 1.
 CC PROSITE; PS00682; ZP_DOMAIN; 1.
 CC Glycoprotein; Signal; Sulfation; Sperm; Receptor; Transmembrane;
 KW Extracellular matrix.
 FT SIGNAL 1 22 POTENTIAL.
 FT CHAIN 23 424 ZONA PELLUCIDA SPERM-BINDING PROTEIN
 FT DOMAIN 23 387 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 388 408 POTENTIAL.
 FT DOMAIN 409 424 CYTOPLASMIC (POTENTIAL).
 FT DOMAIN 45 308 ZP.
 FT CARBOHYD 146 146 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 273 273 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 304 304 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 327 327 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 330 330 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 424 AA; 46303 MW; 9089903FBD268365 CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 424;
 Best Local Similarity 100.0%; Pred. No. 96;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; C
 QY 60 ALGLGLA 66
 Db 389 ALGLGLA 395
 |||||
 |||||
 RESULT 72
 SYH CHLMU
 ID SYH CHLMU STANDARD; PRT; 428 AA.
 AC Q9PUJ9;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Histidyl-tRNA synthetase (EC 6.1.1.21) (Histidine--tRNA ligase)
 DE (Hisers).
 DE HISS OR TC0830.
 OS Chlamydia muridarum.
 OC Bacteria; Chlamydiae; Chlamydiales; Chlamydiaceae; Chlamydia.
 OC NCBI_TaxID=83560;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MoPn / Nigg;
 EX MEDLINE=20150255; PubMed=10684935;
 RA Read T.D., Brunham R.C., Shen C., Gill S.R., Heidelberg J.F.,
 RA White O., Hickey E.K., Peterson J., Utterback T., Berry K., Bass
 RA Linher K., Weidman J., Khouri H., Craven B., Bowman C., Dodson R.
 RA Gwin M., Nelson W., DeBoy R., Kolonay J., McClarty G., Salzberg
 RA Eisen J., Fraser C.M.;
 RA "Genome sequences of Chlamydia trachomatis MoPn and Chlamydia
 RA pneumoniae AR39.";

Res. 28:1397-1406(2000).
 ACTIVITY: ATP + L-histidine + tRNA(His) = AMP +
 e + L-histidyl-tRNA(His).
 omodimer (by similarity).
 R LOCATION: Cytoplasmic.
 : Belongs to class-II aminacyl-tRNA synthetase family.

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 il to license@isb-sib.ch).

 ; AAF39630.1; --
 1QE0.
 -.
 7; -; 1.
 04154; HGTP anticodon.
 04516; His.
 02314; tRNA-synt 2b.
 06195; tRNA-ligase II.
 HGTP anticodon; 1.
 tRNA-synt 2b; 1.
 00442; his; 1.
 62; AA TRNA LIGASE II; 1.
 .synthetase; Protein biosynthesis; Ligase; ATP-binding;
 ome.
 AA; 48939 MW; 9CF859ED0E689DDF CRC64;
 rity 2.5%; Score 7; DB 1; Length 428;
 nservative 100.0%; Pred. No. 96;
 mismatches 0; Mismatches 0; Indels 0; Gaps 0;
 AR 139
 ||
 AR 181
 STANDARD; PRT; 455 AA.
 el. 12, Created)
 el. 12, Last sequence update)
 el. 31, Last annotation update)
 imidine photolysase (EC 4.1.99.3) (DNA photolysase)
 ting enzyme).
 risesus.
 nobacteria; Actinobacteridae; Actinomycetales;
 e; Streptomycetaceae; Streptomycetes.
 1;
 N.A.
 14; PubMed=2501760;
 Takao M., Oikawa A., Yasui A.;
 racterization of a gene encoding a photolysase from
 risesus.";
 Res. 17:4731-4744(1989).
 This enzyme catalyzes the light-dependent monomerization
 (M) of cyclobutyl pyrimidine dimers (in cis-syn
 ion), which are formed between adjacent bases on the
 strand, upon exposure to ultraviolet radiation.
 ACTIVITY: Cyclobutadipyrimidine (in DNA) = 2 pyrimidine
 (in DNA).
 Contains 2 chromophores: a reduced flavin (FADH2) and an
 i-hydroxy-5-deazaflavin (F420). Both chromophores are
 ion-covalent interactions.
 IONS: THIS PROTEIN BELONGS TO THE "LONG WAVELENGTH-TYPE
 IS" WITH AN ABSORPTION MAXIMUM AT ABOUT 440 NM.

CC -!- SIMILARITY: Belongs to the DNA photolysase class-1 family.
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 CC the European Bioinformatics Institute. There are no restrictions
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 CC entities requires a license agreement (See <http://www.isb-sib.ch/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; X15060; CAA33161.1; --
 DR FIR; S05573; S05573.
 DR HSP; P00914; LDNP.
 DR InterPro; IPR002081; DNA photolysase 1.
 DR InterPro; IPR006050; DNA photolysase N.
 DR InterPro; IPR005101; FAD binding 7.
 DR InterPro; IPR006051; FAD binding N.
 DR Pfam; PF00875; DNA photolysase; 1.
 DR Pfam; PF03441; FAD binding 7; 1.
 DR PRINTS; PR00147; DNAPHOTLYASE.
 DR ProDom; PD004390; FAD binding N; 1.
 DR PROSITE; PS00394; DNA PHOTOLYASES 1_1; 1.
 DR PROSITE; PS00691; DNA PHOTOLYASES 1_2; 1.
 KW Lyase; Chromophore; Flavoprotein; FAD; DNA repair; DNA-binding.
 FT DNA BIND 300 319 H-T-H MOTIF (POTENTIAL).
 SQ SEQUENCE 455 AA; 50436 MW; FEA88763DA45931A CRC64;
 Query Match 2.5%; Score 7; DB 1; Length 455;
 Best Local Similarity 100.0%; Pred. No. 1e+02; Indels 0; G
 Matches 7; Conservative 0; Mismatches 0; Indels 0; G
 QY 214 LDLVDG 220
 DB 356 LDLVDG 362
 |||||
 |||||
 RESULT 74
 PUCC RHOC
 ID PUCC RHOC STANDARD; PRT; 461 AA.
 AC P23462;
 DT 01-NOV-1991 (Rel. 20, Created)
 DT 01-NOV-1991 (Rel. 20, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Protein pucC.
 GN PUCC.
 OS Rhodobacter capsulatus (Rhodospseudomonas capsulata).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhodobacterales;
 OC Rhodobacteraceae; Rhodobacter.
 OX NCBI_TaxID=1061;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=89359127; PubMed=2549005;
 RA Tichy H.V., Oberle B., Stiehle H., Schiltz E., Drews G.;
 RT "Genes downstream from pucB and pucA are essential for formation
 RT the B800-850 complex of Rhodobacter capsulatus.";
 RL J. Bacteriol. 171:4914-4922(1989).
 RN [2]
 RP REVISION, AND FUNCTION.
 RX MEDLINE=92007739; PubMed=1717257;
 RA Tichy H.V., Albien K.-U., Gad'On N., Drews G.;
 RT "Analysis of the Rhodobacter capsulatus puc operon: the pucC gene
 RT plays a central role in the regulation of LHII (B800-850 complex)
 RT expression.";
 RL EMBO J. 10:2949-2955(1991).
 RN [3]
 RP TOPOLOGY.
 RX MEDLINE=96326322; PubMed=8759841;
 RA LeBlanc H.N., Beatty J.T.;
 RT "Topological analysis of the Rhodobacter capsulatus PucC protein;
 RT effects of C-terminal deletions on light-harvesting complex II.";
 RL J. Bacteriol. 178:4801-4806(1996).
 CC -!- FUNCTION: PUCC IS REQUIRED FOR HIGH-LEVEL TRANSCRIPTION OF THI
 CC PUC OPERON.

AR LOCATION: Integral membrane protein.
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AA226163.1; ALT_INIT.

C33958.
 004896; PUCC.

ex; Transmembrane.

1 36 CYTOPLASMIC (PROBABLE).
 2 37 PROBABLE.
 3 57 PERIPLASMIC (PROBABLE).
 4 62 PROBABLE.
 5 83 CYTOPLASMIC (PROBABLE).
 6 84 109 CYTOPLASMIC (PROBABLE).
 7 10 129 PROBABLE.
 8 30 142 PERIPLASMIC (PROBABLE).
 9 43 163 PROBABLE.
 10 64 182 CYTOPLASMIC (PROBABLE).
 11 83 202 PROBABLE.
 12 103 209 PERIPLASMIC (PROBABLE).
 13 110 228 PROBABLE.
 14 129 261 CYTOPLASMIC (PROBABLE).
 15 162 281 PROBABLE.
 16 192 302 PERIPLASMIC (PROBABLE).
 17 103 319 PROBABLE.
 18 120 338 CYTOPLASMIC (PROBABLE).
 19 139 355 PROBABLE.
 20 156 358 PERIPLASMIC (PROBABLE).
 21 159 376 PROBABLE.
 22 177 394 CYTOPLASMIC (PROBABLE).
 23 195 415 PROBABLE.
 24 116 436 PERIPLASMIC (PROBABLE).
 25 137 456 PROBABLE.
 26 157 461 CYTOPLASMIC (PROBABLE).
 27 11 AA; 48392 MW; BFC7A8A0C549875A CRC64;

2.5%; Score 7; DB 1; Length 461;
 arity 100.0%; Pred. No. 1e+02;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;

EDQ 101

EDQ 176

STANDARD; PRT; 461 AA.

(Rel. 35, Created)
 (Rel. 35, Last sequence update)
 (Rel. 41, Last annotation update)
 protein HI0608.

influenzae.
 teobacteria; Gammaproteobacteria; Pasteurellales;
 ae; Haemophilus.

4 N.A.

0630 / ATCC 51907;

0630; PubMed=7542800;

3.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
 3., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
 Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
 Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
 Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,

RA Utterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.
 RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
 RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
 RA Venter J.C.;
 RT "Whole-genome random sequencing and assembly of Haemophilus infl
 RT Rd.";
 RL Science 269:496-512(1995).
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
 CC -!- SIMILARITY: BELONGS TO THE SLC13A FAMILY OF TRANSPORTERS.
 CC NADC SUBFAMILY.

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 CC between the Swiss Institute of Bioinformatics and the EMBL out
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CC EMBL; U32743; AAC22267.1; -.

DR PIR; I64080; I64080.

DR TIGR; HI0608; -.

DR InterPro; IPR001898; Na/sul_sympo.

DR Pfam; PF00939; Na_sulph_symp; 1.

DR TIGRFAMs; TIGR00785; dars; 1.

DR PROSITE; PS01271; NA_SULFATE; 1.

KW Hypothetical protein; Transmembrane; Transport; Complete proteom
 FT TRANSMEM 13 33 POTENTIAL.
 FT TRANSMEM 54 74 POTENTIAL.
 FT TRANSMEM 81 101 POTENTIAL.
 FT TRANSMEM 120 140 POTENTIAL.
 FT TRANSMEM 170 190 POTENTIAL.
 FT TRANSMEM 211 231 POTENTIAL.
 FT TRANSMEM 256 276 POTENTIAL.
 FT TRANSMEM 286 306 POTENTIAL.
 FT TRANSMEM 314 334 POTENTIAL.
 FT TRANSMEM 349 369 POTENTIAL.
 FT TRANSMEM 377 397 POTENTIAL.
 FT TRANSMEM 399 419 POTENTIAL.
 FT TRANSMEM 439 459 POTENTIAL.
 SQ SEQUENCE 461 AA; 49761 MW; B5E6F6965B38EF06 CRC64;

Query Match

2.5%; Score 7; DB 1; Length 461;

Best Local Similarity 100.0%; Pred. No. 1e+02;

Matches 7; Conservative 0; Mismatches 0; Indels 0;

QY 53 TALLVPL 59

Db 60 TALLVPL 66

Search completed: April 7, 2004, 17:57:56

Job time : 21 secs

16:25:21 2004

us-09-245-198a-4.oligo.rspt

GenCore version 5.1.6
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n search, using sw model

il 7, 2004, 17:54:13 ; Search time 45 Seconds
(without alignments)
1991.270 Million cell updates/sec

09-245-198A-4

MSLLDFEISARRLPRLSLG.....PWAHLKAAPELTIVGLFQVH 284

GO

xop 60.0 , Gapext 60.0

.7041 seqs, 315518202 residues

s satisfying chosen parameters: 1017041

jth: 0

jth: 2000000000

sting first 100 summaries

PREMBL 25:*

sp archaea:*

sp bacteria:*

sp fungi:*

sp human:*

sp invertebrate:*

sp mammal:*

sp_mhc:*

sp_organelle:*

sp_phage:*

sp_plant:*

sp_rodent:*

sp_virus:*

sp_vertebrate:*

sp_unclassified:*

sp_rvirus:*

sp_bacteriap:*

sp_archaeap:*

the number of results predicted by chance to have a
r than or equal to the score of the result being printed,
ad by analysis of the total score distribution.

SUMMARIES

ary	Length	DB	ID	Description
3.5	330	4	Q8IZK7	Q8izk7 homo sapien
1.3	410	11	Q8BXS2	Q8bxs2 mus musculus
4.2	438	16	Q7VVB7	Q7vvb7 bordetella
4.2	470	16	Q7W7P2	Q7wf2 bordetella
3.5	111	16	Q8XA78	Q8x478 escherichia
3.5	111	16	Q8PFL8	Q8ffl8 escherichia
3.5	111	16	Q7UC61	Q7uc61 shigella fl
3.2	142	16	Q8BIJ0	Q8bi0 rhizobium l
3.2	749	16	Q7V511	Q7v511 prochloroco
3.2	766	16	Q8PZ8	Q8ppz8 xanthomonas
3.2	1208	16	Q7ULK4	Q7ulk4 rhodopirell
2.8	143	17	Q9HST7	Q9hst7 halobacteri
2.8	151	10	Q9SD11	Q9sdl1 oryza sativ
2.8	154	2	Q848K4	Q848k4 gamma-prote
2.8	190	16	Q8E569	Q8e569 streptococc
2.8	190	16	Q8DZK5	Q8dzk5 streptococc

Q8L4k2 ory
Q9wv10 the
Q9vv70 dros
Q8sxh4 dros
Q8fv59 bru
Q89w9 bra
Q84mb7 ara
Q9z999 pse
Q82143 str
Q9d378 mus
Q9cpr8 mus
Q9vnp0 dros
Q86411 pan
Q86410 pong
Q86418 maca
Q86417 maca
Q86415 maca
Q86498 actu
Q86497 atel
Q86496 alou
Q86494 alou
Q86493 alou
Q86492 alou
Q86491 alou
Q86418 vare
Q86417 vare
Q86416 eule
Q86415 hapa
Q86414 lemu
Q85554 pyr
Q99908 homo
Q9bpv2 homo
Q8tuu8 met
Q53860 myc
Q7u145 myc
Q877C5 meth
Q9kyt3 str
Q9shd8 ara
Q8u820 agr
Q7u9h4 syn
Q8xs70 ral
Q8zc8 yer
Q9rr18 dei
Q96n66 homo
Q8rip9 mus
Q9cy76 mus
Q8chk3 mus
Q8gvz8 ory
Q7wz71 nono
Q8pmh8 xan
Q9vtu9 inf
Q8j192 inf
Q8qpl0 inf
Q8pas2 xan
Q9n8h2 tryp
Q9n8u8 tryp
Q9bef6 capr
Q9z935 chla
Q16193 homo
Q7u694 syn
Q8rq64 serr
Q8h142 ara
Q9a4q4 cau
Q9p074 homo
Q82cb6 nit
Q7x682 ory
P97199 esch
Q8qtr8 eupr
Q8qrh6 hep
Q8utl1 hum
Q8dbe3 vib
Q38214 bact

2.5 115 16 Q9HY60
 2.5 118 2 Q939F7
 2.5 118 16 Q928G5
 2.5 119 10 Q94LX6
 2.5 121 10 Q8W0M1
 2.5 122 10 Q84PV3
 2.5 123 16 Q8EFG2
 2.5 123 16 Q8UGH2
 2.5 123 16 Q7U823
 2.5 124 10 Q7XRK7
 2.5 125 5 Q9U1P6

ALIGNMENTS

PRELIMINARY; PRT; 330 AA.

TrEMBLrel. 23, Created
 TrEMBLrel. 23, Last sequence update
 TrEMBLrel. 25, Last annotation update

(Human).
 Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Primates; Catarrhini; Hominidae; Homo.
 106;

1 N.A.
 1924; PubMed=12411489;
 B.; Medema J.P.; Lopez-Fraga M.; Lozano J.C.,
 M.; Picard A., Martinez A.C., Garcia-Sanz J.A.,

is hybrid mRNA encodes TWE-PRIL, a functional cell surface
 fusion protein."
 11-5720(2002).

1; AAL90443.1; -
 0; C.membrane; IEA.
 4; F.tumor necrosis factor receptor binding; IEA.
 5; P.immune response; IEA.
 106052; TNF family.
 1008983; TNF_like.
 1; TNF; 1.
 17; TNF; 1.
 1251; TNF 1; 1.
 1049; TNF 2; 2.
 10 AA; 36588 MW; FC6F3BCA29C029AE CRC64;

58.5%; Score 166; DB 4; Length 330;
 arity 100.0%; Pred. No. 1.3e-155;
 conservative 0; Mismatches 0; Indels 0; Gaps 0;

RSQRGRGEGPTALLVPLALGLALACGLLLAVVSLGSRASLSAQEAQEL 95
 RSQRGRGEGPTALLVPLALGLALACGLLLAVVSLGSRASLSAQEAQEL 60
 RDPSFLNPQTEESQDPAFLNRLVPRSPKGRKTRRAIAAHYVHPRPGD 155
 RDPSFLNPQTEESQDPAFLNRLVPRSPKGRKTRRAIAAHYVHPRPGD 120

AGVDGTVSGWEARINSSPLRNQIGFIVTRAGLYLYYCQ 201
 AGVDGTVSGWEARINSSPLRNQIGFIVTRAGLYLYYCQ 166

PRELIMINARY; PRT; 410 AA.

(TrEMBLrel. 23, Created)
 (TrEMBLrel. 23, Last sequence update)
 (TrEMBLrel. 25, Last annotation update)

DE Tumor necrosis factor.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus
 OC NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Retina;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium,
 RA the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotati
 RT 60,770 full-length cDNAs.";
 RL Nature 420:563-573(2002).
 DR EMBL; AK044387; BAC31897.1; -
 DR PIR; PT0714; PT0714.
 DR GO; GO:0016020; C.membrane; IEA.
 DR GO; GO:0005184; F.tumor necrosis factor receptor binding; IEA.
 DR GO; GO:0006955; P.immune response; IEA.
 DR InterPro; IPR006052; TNF family.
 DR InterPro; IPR008983; TNF_like.
 DR SMART; SM00207; TNF; 2.
 DR PROSITE; PS00251; TNF 1; 1.
 DR PROSITE; PS0049; TNF 2; 2.
 SQ SEQUENCE 410 AA; 45881 MW; 590A4B74C33FB8D4 CRC64;

Query Match 11.3%; Score 32; DB 11; Length 410;
 Best Local Similarity 100.0%; Pred. No. 8.9e-23;
 Matches 32; Conservative 0; Mismatches 0; Indels 0; C

QY 139 RRAIAAHYVHPRPGQDGAQAGVDGTVSGWEE 170
 DB 105 RRAIAAHYVHPRPGQDGAQAGVDGTVSGWEE 136

RESULT 3

Q7VVB7
 ID Q7VVB7 PRELIMINARY; PRT; 438 AA.
 AC Q7VVB7;
 DT 01-OCT-2003 (TrEMBLrel. 25, Created)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Putative chloride-channel protein.
 GN BP2760.
 OS Bordetella pertussis.
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
 OC Alcaligenaceae; Bordetella.
 OC NCBI_TaxID=520;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Tohama I / ATCC BAA-589 / NCTC 13251;
 RX MEDLINE=22827954; PubMed=12910271;
 RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,
 RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall
 RA Cerdeno-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,
 RA Achtman M., Ackin R., Baker S., Basham D., Bason N., Cherevach I.
 RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,
 RA Felwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels I
 RA Leather S., Moule S., Norberczak H., O'Neil S., Ormond D., Price
 RA Rabinowitch E., Rutter S., Sanders M., Saunders D., Seeger K.,
 RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Steve
 RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;
 RT "Comparative analysis of the genome sequences of Bordetella pert
 RT Bordetella parapertussis and Bordetella bronchiseptica.";
 RL Nat. Genet. 35:32-40(2003).
 DR EMBL; EX640419; CAE43035.1; -
 KW Complete proteome.
 SQ SEQUENCE 438 AA; 45402 MW; AD51A65B59599D8 CRC64;

Query Match 4.2%; Score 12; DB 16; Length 438;
 Best Local Similarity 100.0%; Pred. No. 0.0062;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; C

ALACLGLL 73
|||||
ALACLGLL 285

RELIMINARY; PRT; 470 AA.

TrEMBLrel. 25, Created)
TrEMBLrel. 25, Last sequence update)
TrEMBLrel. 25, Last annotation update)
side-channel protein.

apertussis.
eobacteria; Betaproteobacteria; Burkholderiales;
; Bordetella.

N.A.

/ ATCC BAA-587;
954; PubMed=12910271;
Bebahia M., Preston A., Murphy L.D., Thomson N.,
olden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,
A.M., Temple L., James K., Harris B., Quail M.A.,
kin R., Baker S., Basham D., Bason N., Cherevach I.,
T., Collins M., Cronin A., Davis P., Doggett J.,
oble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,
ule S., Norberczak H., O'Neill S., Ormond D., Price C.,
E., Rutter S., Sanders M., Saunders D., Seeger K.,
onds M., Skelton J., Squares R., Squares S., Stevens K.,
head S., Barrell B.G., Maskell D.J.;
analysis of the genome sequences of Bordetella pertussis,
apertussis and Bordetella bronchiseptica.";
3:32-40(2003).
); CAB37862.1; -.
); AA; 49233 MW; 5B92DD05E920BDSF CRC64;

4.2%; Score 12; DB 16; Length 470;

arity 100.0%; Pred. No. 0.066;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

ALACLGLL 73
|||||
ALACLGLL 317

RELIMINARY; PRT; 111 AA.

TrEMBLrel. 20, Created)
TrEMBLrel. 20, Last sequence update)
TrEMBLrel. 24, Last annotation update)
protein z3516.

oli O157:H7.
eobacteria; Gammaproteobacteria; Enterobacteriales;
334;
Escherichia.

N.A.

7 / EDL933 / ATCC 700927;
935; PubMed=11206551;
lunkett G. III, Burland V., Mau B., Glauner J.D.,
ynew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,
ckert J., Klink S., Boutin A., Shao Y., Miller L.,
Davis N.W., Lim A., Dimalanta E.T., Potamocis K.,
nantharaman T.S., Lin J., Yen G., Schwartz D.C.,
lattner F.R.;
nce of enterohaemorrhagic Escherichia coli O157:H7.";
9-533(2001).

DR EMBL; AB005458; AAG57389.1; -.
DR PIR; A85866; A85866.
DR GO; GO:0016020; C:membrane; IEA.
DR InterPro; IPR000620; DUF6.
DR Pfam; PF00892; DUF6; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 111 AA; 12165 MW; 7CEFC93D786CD759 CRC64;

Query Match 3.5%; Score 10; DB 16; Length 111;
Best Local Similarity 100.0%; Pred. No. 0.17;
Matches 10; Conservative 0; Mismatches 0; Indels 0; G

QY 63 LGLALACLG 72
|||||
DB 40 LGLALACLG 49

RESULT 6

Q8FFL8 PRELIMINARY; PRT; 111 AA.
ID Q8FFL8
AC Q8FFL8
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Conserved hypothetical protein.
GN C2800.
OS Escherichia coli O6.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=217992;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=06:H1 / CFT073 / ATCC 700928;
RX MEDLINE=22388234; PubMed=12471157;
RA Weich R.A., Burland V., Plunkett G. III, Redford P., Roesch P.,
RA Rasko D., Buckles E.L., Liou S.-R., Boutin A., Hackett J., Stroud
RA Mayhew G.F., Rose D.J., Zhou S., Schwartz D.C., Perna N.T.,
RA Mobley H.L.T., Donnenberg M.S., Blattner F.R.;
RT "Extensive mosaic structure revealed by the complete genome seque
of uropathogenic Escherichia coli".
RL Proc. Natl. Acad. Sci. U.S.A. 99:17020-17024(2002).
DR EMBL; AE016763; AN81254.1; -.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 111 AA; 12196 MW; C0A977B6F77A4B87 CRC64;

Query Match 3.5%; Score 10; DB 16; Length 111;
Best Local Similarity 100.0%; Pred. No. 0.17;
Matches 10; Conservative 0; Mismatches 0; Indels 0; G

QY 63 LGLALACLG 72
|||||
DB 40 LGLALACLG 49

RESULT 7

Q7UC61 PRELIMINARY; PRT; 111 AA.
ID Q7UC61
AC Q7UC61
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Sucrose-6 phosphate hydrolase.
GN S2567.
OS Shigella flexneri.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Shigella.
OX NCBI_TaxID=623;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=2457T / ATCC 700930 / Serotype 2a;
RX MEDLINE=22590274; PubMed=12704152;
RA Wei J., Goldberg M.B., Burland V., Venkatesan M.M., Deng W.,
RA Fournier G., Mayhew G.F., Plunkett G. III, Rose D.J., Darling A.,

1 N.T., Payne S.M., Runyen-Janecky L.J., Zhou S.,
Blattner F.R.;
ome sequence and comparative genomics of Shigella
type 2a strain 2457T";
1. 71:2775-2786(2003).
36; AAP17670.1; -.

11 AA; 12224 MW; 7CFA06CC46A32672 CRC64;

3.5%; Score 10; DB 16; Length 111;
larity 100.0%; Pred. No. 0.17; 0; Indels 0; Gaps 0;
Conservative 0; Mismatches 0; Mismatches 0; Indels 0;

ALACGL 72

|||||

ALACGL 49

PRELIMINARY; PRT; 142 AA.

(TREMBLrel. 18, Created)

(TREMBLrel. 18, Last sequence update)

(TREMBLrel. 18, Last annotation update)

nsport protein.

ti (Mesorhizobium loti).

teobacteria; Alphaproteobacteria; Rhizobiales;

iaceae; Mesorhizobium.

91;

M N.A.

03099;

2930; PubMed=11214968;

akamura Y., Sato S., Asamizu E., Kato T., Sasamoto S.,

Ideawa K., Ishikawa A., Kawashima K., Kimura T.,

Kiyokawa C., Kohara M., Matsumoto M., Matsuno A.,

Nakayama S., Nakazaki N., Shimpo S., Sugimoto M.,

Yanada M., Tabata S.;

name structure of the nitrogen-fixing symbiotic bacterium

m loti";

31-338(2000).

99; BAB49526.1; -.

teone.

42 AA; 14884 MW; ODCA7842C85A5B6F CRC64;

3.2%; Score 9; DB 16; Length 142;

larity 100.0%; Pred. No. 2.1; 0; Indels 0; Gaps 0;

Conservative 0; Mismatches 0; Mismatches 0; Indels 0;

LVPLAL 61

|||||

LVPLAL 130

PRELIMINARY; PRT; 749 AA.

(TREMBLrel. 25, Created)

(TREMBLrel. 25, Last sequence update)

(TREMBLrel. 25, Last annotation update)

I PeaB protein.

769.

cus marinus (strain MIT 9313).

anobacteria; Prochlorophytes; Prochlorococcaceae;

cus.

4547;

M N.A.

5698; PubMed=12917642;

zimer F.W., Lamerdin J., Malfatti S., Chain P.,

RA Ahlgren N.A., Arellano A., Coleman M., Hauser L., Hess W.R.,
RA Johnson Z.I., Land M., Lindell D., Post A.F., Regalia W., Shah M.
RA Shaw S.L., Steglich C., Sullivan M.B., Ting C.S., Tolonen A.,
RA Webb E.A., Zinser E.R., Chisholm S.W.;

"Genome divergence in two Prochlorococcus ecotypes reflects ocea

RT niche differentiation";

RL Nature 424:1042-1047(2003).

DR EMBL; BX572100; CAE21944.1; -.

KW Photosystem I; Complete proteome.

SQ SEQUENCE 749 AA; 83231 MW; B1D496645F1C790C CRC64;

Query Match 3.2%; Score 9; DB 16; Length 749;

Best Local Similarity 100.0%; Pred. No. 9.4; 0; Indels 0;

Matches 9; Conservative 0; Mismatches 0; Mismatches 0;

QY 63 LGLALACLG 71

|||||

Db 347 LGLALACLG 355

RESULT 10

Q8PPZ8

ID Q8PPZ8 PRELIMINARY; PRT; 766 AA.

AC Q8PPZ8;

DT 01-OCT-2002 (TREMBLrel. 22, Created)

DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)

DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)

DE C-type cytochrome biogenesis protein (Copper tolerance).

GN DSBD OR XAC0534.

OS Xanthomonas axonopodis (pv. citri).

OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;

OC Xanthomonadaceae; Xanthomonas.

OX NCBI_TaxID=92829;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=306 / ATCC 13902 / XV 101;

RE MEDLINE=22022145; PubMed=12024217;

RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.

RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.

RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,

RA Camarotte G., Cannavaro F., Cardoso J., Chambergo F., Clapina L.P.

RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorri H.

RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Gruber A.,

RA Formighieri E.F., Franco M.C., Greggio C.C., Lemos M.V.

RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.

RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.

RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,

RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.

RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.

RA Spinoia L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.

RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,

RA Setubal J.C., Kitajima J.P.;

RT "Comparison of the genomes of two Xanthomonas pathogens with dif

RT host specificities";

RL Nature 417:459-463(2002).

DR EMBL; AF011680; AM35423.1; -.

DR GO; GO:0016020; C:membrane; IEA.

DR GO; GO:0005489; F:electron transporter activity; IEA.

DR GO; GO:0017004; P:cytochrome biogenesis; IEA.

DR GO; GO:0006118; P:electron transport; IEA.

DR InterPro; IPR003834; Cytococh.TM.

DR InterPro; IPR006662; ThioRed.

DR InterPro; IPR006663; ThioRedox_dom2.

DR Pfam; PF02683; DSbd; 1.

DR PROSITE; PS00194; THIOREDOXIN; 1.

KW Complete proteome.

SQ SEQUENCE 766 AA; 81014 MW; 3A1955A07DB8A9CA CRC64;

Query Match 3.2%; Score 9; DB 16; Length 766;

Best Local Similarity 100.0%; Pred. No. 9.6;

Matches 9; Conservative 0; Mismatches 0; Indels 0;

QY 59 LAIGLGLAL 67

|||||
GLAL 542

RELIMINARY; PRT; 1208 AA.

TrEMBLrel. 25, Created)
TrEMBLrel. 25, Last sequence update)
TrEMBLrel. 25, Last annotation update)
; multi-functional protein.

; baltica.
; ctomycetes; Planctomycetacia; Planctomycetales;
; eae; Pirellula.
; ;

N.A.

113; PubMed=12835416;
; Kube M., Bauer M., Teeling H., Lombardot T.,
; ie D., Beck A., Borzym K., Heilmann K., Rabus R.,
; Anann R., Reinhardt R.;
; me sequence of the marine planctomycete Pirellula sp.

ad. Sci. U.S.A. 100:8298-8303 (2003).

; CAD76265.1; -.

18 AA; 132047 MW; OFFE225741021E8C CRC64;

3.2%; Score 9; DB 16; Length 1208;

arity 100.0%; Pred. No. 15;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

AOEPA 91

|||||

AOEPA 42

RELIMINARY; PRT; 143 AA.

TrEMBLrel. 16, Created)
TrEMBLrel. 16, Last sequence update)
TrEMBLrel. 24, Last annotation update)

sp. (strain NRC-1 / ATCC 700922 / JCM 11081).

archaeota; Halobacteria; Halobacteriales;

ae; Halobacterium.

391;

N.A.

483; PubMed=11016950;
edy S.P., Mahairas G.G., Berquist B., Pan M.,
asky S.R., Balliga N.S., Thorsson V., Shrogha J.,
Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,
; Keller K., Cruz R., Danson M.J., Hough D.W.,
; Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,
A., Peck R.F., Pohlchröder M., Spudich J.L., Jung K.-H.,
tas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,
owe T.M., Liang P., Riley M., Hood L., Dassarma S.;
nce of Halobacterium species NRC-1.;

cad. Sci. U.S.A. 97:12176-12181 (2000).

6; AAG18715.1; -.

G84168.

006976; Vanz.

; Vanz; 1.

some.

3 AA; 15648 MW; 45466B6328EF3468 CRC64;

Query Match 2.8%; Score 8; DB 17; Length 143;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 61 LGLGLALA 68
Db 55 LGLGLALA 62

RESULT 13

Q9SD11

ID Q9SD11 PRELIMINARY; PRT; 151 AA.

AC Q9SD11;

DT 01-MAY-2000 (TrEMBLrel. 13, Created)

DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)

DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)

DE Hypothetical protein (OSUNBA0036E02.6 protein) (B1085F09.2

DE protein).

GN B1085F09.2.

OS Oryza sativa (Rice).

OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta

OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

OC Ehrhartoideae; Oryzeae; Oryza.

OX NCBI_TaxID=4530;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=cv. Nipponbare;

RA Sasaki T., Matsumoto T., Yamamoto K.;

RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, PAC

clone:P0003H10.1; -;

RL Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=cv. Nipponbare;

RA Sasaki T., Matsumoto T., Yamamoto K.;

RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, BAC

clone:OSUNBA0036E02.1; -;

RL Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.

RN [3]

RP SEQUENCE FROM N.A.

RC STRAIN=cv. Nipponbare;

RA Sasaki T., Matsumoto T., Yamamoto K.;

RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, BAC

clone:B1085F09.1; -;

RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.

DR EMBL; AP000815; BAA87834.1; -.

DR EMBL; AP002862; BAB17732.1; -.

DR EMBL; AP003103; BAB44106.1; -.

DR Gramene; Q9SD11; -.

SQ SEQUENCE 151 AA; 16632 MW; EC68451ECA2BD71D CRC64;

Query Match 2.8%; Score 8; DB 10; Length 151;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 43 RRRGRGGE 50
Db 131 RRRGRGGE 138

RESULT 14

Q848K4

ID Q848K4 PRELIMINARY; PRT; 154 AA.

AC Q848K4;

DT 01-JUN-2003 (TrEMBLrel. 24, Created)

DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)

DE Hypothetical protein (Fragment).

OS Gamma-proteobacterium Hot 75m4.

OG Plasmid pAK106.

OC Bacteria; environmental samples.

OX NCBI_TaxID=77133;

RN [1]

```

1 N.A.
7661; PubMed=12620823;
Waschkowitz T., Bowien S., Henne A., Daniel R.;
1 and Screening of Metagenomic Libraries Derived from
1ltures: Generation of a Gene Bank for Genes Conferring
reductase Activity on Escherichia coli.";
1. Microbiol. 69:1408-1416(2003).
21; AAO59972.1; -.
21; C:extrachromosomal DNA; IEA.
protein; Plasmid.
1
54 AA; 16234 MW; 3AE8A072D5B7E137 CRC64;
arity 2.8%; Score 8; DB 2; Length 154;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
}GRRG 49
|||||
}GRRG 119

PRELIMINARY; PRT; 190 AA.
(TREMBLrel. 23, Created)
(TREMBLrel. 23, Last sequence update)
(TREMBLrel. 24, Last annotation update)
protein.

s agalactiae (serotype III).
rmicutes; Lactobacillales; Streptococcaceae;
s.
16495;

N.A.
5 / Serotype III;
2508; PubMed=12354221;
usniok C., Buchrieser C., Chevallier F., Frangeul L.,
cuine M., Couve E., Lalioui L., Poyart C., Trieu-Cuot P.,
ence of Streptococcus agalactiae, a pathogen causing
natal disease.";
21. 45:1499-1513(2002).
49; CAB46822.1; -.
s1163; -.
R008172; Adenylate_cyc.
8; CYTH; 1.
protein; Complete proteome.
90 AA; 22178 MW; AB2AD33C2CB6FBF3 CRC64;
arity 2.8%; Score 8; DB 16; Length 190;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

IRITLP 265
|||||
IRITLP 57

PRELIMINARY; PRT; 190 AA.
(TREMBLrel. 23, Created)
(TREMBLrel. 23, Last sequence update)
(TREMBLrel. 24, Last annotation update)
pothetical protein.

s agalactiae (serotype V).
rmicutes; Lactobacillales; Streptococcaceae;
s.

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OX NCBI_TaxID=216466;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=2603 V/R / Serotype V;
RC MEDLINE=22222988; PubMed=12200547;
RA Tettelin H., Masignani V., Cieslewicz M.J., Eisen J.A., Peterson
RA Wessels M.R., Paulsen I.T., Nelson K.E., Margarit I., Read T.D.,
RA Madoff L.C., Wolf A.M., Beanan M.J., Brinkac L.M., Daugherty S.C
RA DeBoy R.T., Durkin A.S., Klonay J.F., Madupu R., Lewis M.R.,
RA Radune D., Fedorova N.B., Scanlan D., Khouri H., Mulligan S.,
RA Carthy H.A., Cline R.T., Van Aken S.E., Gill J., Scarselli M., Mo:
RA Iacobini E.T., Brettoni C., Galli G., Mariani M., Vegni F., Maior
RA Rinaldo D., Rappuoli R., Telford J.L., Kasper D.L., Grandi G.,
RA Fraser C.M.;
RT "Complete genome sequence and comparative genomic analysis of an
RT emerging human pathogen, serotype V Streptococcus agalactiae.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:12391-12396(2002).
DR EMBL; AE014242; AAM99977.1; -.
DR TIGR; SAG1096; -.
DR InterPro; IPR008172; Adenylate_cyc.
DR Pfam; PF01928; CYTH; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 190 AA; 22178 MW; AB2AD33C2CB6FBF3 CRC64;

Query Match 2.8%; Score 8; DB 16; Length 190;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 258 SLRIRTLP 265
Db |||||
50 SLRIRTLP 57

RESULT 17
Q8L4K2 PRELIMINARY; PRT; 193 AA.
AC Q8L4K2;
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN OSJNBA0079H13.8 OR OSJNBA0038H12.21.
OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyt
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzaceae; Oryza.
OX NCBI_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA Buell C.R., Yuan Q., Ouyang S., Liu J., Gansberger K., Kim M.M.,
RA Overton II L.L., Bera J.J., Tsitrin T., Krol M.I., Jarrahi B.B.,
RA Jin S.S., Koo H., Zismann V., Hsiao J., Blunt S., Vanaken S.S.,
RA Utterback T.T., Feldblyum T.V., Yang Q.Q., Haas B.J., Suh B.B.,
RA Peterson J.J., Quackenbush J., White O., Salzberg S.L., Fraser C
RA "Oryza sativa chromosome 10 BAC OSJNBA0079H13 genomic sequence."
RT Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
RL [2]
RN SEQUENCE FROM N.A.
RP STRAIN=cv. Nipponbare;
RA Buell C.R., Yuan Q., Ouyang S., Liu J., Gansberger K., Jones K.M
RA Overton II L.L., Tsitrin T., Kim M.M., Bera J.J., Jin S.S.
RA Fadrosch D.W., Tallon L.J., Koo H., Zismann V., Hsiao J., Blunt S
RA Vanaken S.S., Riedmuller S.B., Utterback T.T., Feldblyum T.V.,
RA Yang Q.Q., Haas B.J., Suh B.B., Peterson J.J., Quackenbush J.,
RA White O., Salzberg S.L., Fraser C.M.;
RA "Oryza sativa chromosome 10 BAC OSJNBA0038H12 genomic sequence."
RT Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
RL [3]
RN SEQUENCE FROM N.A.
RP STRAIN=cv. Nipponbare;
RC The Rice Chromosome 10 Sequencing Consortium;
RA "In-depth view of structure, activity, and evolution of rice

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";
66-1569 (2003).
N.A.
ponbare;
ng R.A., McCombie W.R., Messing J., Yuan Q.;
-2003) to the EMBL/GenBank/DBJ databases.
; AAM54153.1; -
; AAN04965.1; -
; AAP52530.1; -
2; -
rotein.
AA; 20812 MW; 719544BFC9A0790 CRC64;
2.8%; Score 8; DB 10; Length 193;
urity 100.0%; Pred. No. 27; Indels 0; Gaps 0;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
IRGE 50
|||||
RGE 123

ELIMINARY; PRT; 197 AA.

'EMBLrel. 12, Created)
'EMBLrel. 12, Last sequence update)
'EMBLrel. 24, Last annotation update)
rotein TW0469.

itima
motogae; Thermotogales; Thermotogaceae; Thermotoga.
16;
N.A.
DSM 3109;
115; PubMed:10360571;
layton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
Key E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
Jeterback T.R., Malek J.A., Linher K.D., Garrett M.M.,
Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
Smith H.O., Venter J.C., Fraser C.M.;
lateral gene transfer between Archaea and Bacteria from
e of Thermotoga maritima.";
1-329(1999).
; AAD35553.1; -
; 272374.
rotein; Complete proteome.
/ AA; 22919 MW; 41E2C8E3C09180EC CRC64;
2.8%; Score 8; DB 16; Length 197;
urity 100.0%; Pred. No. 28; Indels 0; Gaps 0;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
EIS 9
|||||
EIS 142

ELIMINARY; PRT; 211 AA.

'EMBLrel. 13, Created)
'EMBLrel. 22, Last sequence update)
'EMBLrel. 24, Last annotation update)
in.
lanogaster (Fruit fly).

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```

OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
RN [1]
SEQUENCE FROM N.A.
RC STRAIN=Berkeley;
RX MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N
RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L
RA Abril J.F., Aghayani A., An H.-J., Andrews-Pfannkuch C., Baldwin I
RA Ballaw R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M
RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,
RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dum
RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischman
RA Fostler C., Gabrielian A.B., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwac C.,
RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum I
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasarman D.A., Weinstock G.M., Weisenbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2195-2195 (2000).
RN [2]
SEQUENCE FROM N.A.
RP Celniker S.E., Adams M.D., Krommiller B., Wan K.H., Holt R.A.,
RA Evans C.A., Gocayne J.D., Amanatides P.G., Brandon R.C., Rogers Y
RA Banzon J., An H., Baldwin D., Banzon J., Beeson K.Y., Busam D.A.,
RA Carlson J.W., Center A., Champe M., Davenport L.B., Dietz S.M.,
RA Dedson K., Dorsett V., Doup L.E., Doyle C., Dresnek D., Farfan D.
RA Ferreira S., Frise E., Galie R.F., Garg N.S., George R.A.,
RA Gonzalez C., Houck J., Hoskins R.A., Hostin D., Howland T.J.,
RA Ibegwac M., Jallali M., Kruse D., Li P., Mattei B., Moshrefi A.,
RA McIntosh T.C., Moy M., Murphy B., Nelson C., Nelson K.A., Nunco J
RA Pacleb J., Paragas V., Park S., Patel S., Pfeiffer B.,
RA Prounenavong S., Pittman G.S., Puri V., Richards S., Scheeler F.
RA Stapleton M., Strong R., Svirskas R., Tector C., Tyler D.,
RA Williams S.M., Zaveri J.S., Smith H.O., Venter J.C., Rubin G.M.;
RT "Sequencing of Drosophila melanogaster genome.";
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
SEQUENCE FROM N.A.
RP Misra S., Crosby M.A., Matthews B.B., Bayraktaroglu L., Campbell
RA Hradscky P., Huang Y., Kaminker J.S., Prochink S.E., Smith C.D.,
RA Tupy J.L., Bergman C., Berman B., Carlson J.W., Celniker S.E.,
RA Clamp M., Drysdale R., Emert D., Frise E., de Grey A., Harris N.
RA Krommiller B., Marshall B., Millburn G., Richter J., Russo S.,
RA Searle S.M.J., Smith E., Shu S., Smutniak F., Whitfield E.,
RA Ashburner M., Gelbart W.M., Rubin G.M., Mungall C.J., Lewis S.E.;
RT "Annotation of Drosophila melanogaster genome.";

```

R-2000) to the EMBL/GenBank/DBJ databases.

(N.A.
elinker S.E., Gibbs R.A., Rubin G.M., Venter C.J.;
R-2000) to the EMBL/GenBank/DBJ databases.

(N.A.

P-2002) to the EMBL/GenBank/DBJ databases.

7; RAF49452.2; -

0036638; CGI3033.

004011; Gyr.

004019; YLP_motif.

; Gyr; 1.

; YLP; 5.

1 AA; 23779 MW; D2554983E91F5107 CRC64;

2.8%; Score 8; DB 5; Length 211;
arity 100.0%; Pred. No. 29;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

AVVSL 79

|||||

AVVSL 16

PRELIMINARY; PRT; 211 AA.

TrEMBLrel. 21, Created)

TrEMBLrel. 21, Last sequence update)

TrEMBLrel. 24, Last annotation update)

Planogaster (Fruit fly).

stazoa; Arthropoda; Hexapoda; Insecta; Pterygota;

lopterygota; Diptera; Brachycera; Muscomorpha;

Drosophilidae; Drosophila.

27;

(N.A.

ey;

Brokstein P., Hong L., Agbayani A., Carlson J.,

avez C., Dorsett V., Dresnek D., Farfan D., Friese E.,

nzalez M., Guarin H., Krommiller B., Li P., Liao G.,

ngall C.J., Nunco J., Pacleb J., Faragas V., Park S.,

uanenavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,

AR-2002) to the EMBL/GenBank/DBJ databases.

4; AAL90372.1; -

0063673; BCDNA.RE50345.

004011; Gyr.

004019; YLP_motif.

; Gyr; 1.

; YLP; 5.

11 AA; 23780 MW; 82FF4983E91F510A CRC64;

2.8%; Score 8; DB 5; Length 211;
arity 100.0%; Pred. No. 29;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

AVVSL 79

|||||

AVVSL 16

PRELIMINARY; PRT; 220 AA.

(TrEMBLrel. 23, Created)

(TrEMBLrel. 23, Last sequence update)

DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)

DE Membrane protein, putative.

GN BRA0991.

OS Brucella suis.

OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;

OC Brucellaceae; Brucella.

OX NCBI_TaxID=29461;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=1330 / Biovar 1;

RX MEDLINE=22247741; PubMed=12271122;

RA Paulsen I.T., Seeshadri R., Nelson K.E., Eisen J.A., Heidelberg J.

RA Read T.D., Dodson R.J., Unayam L., Brinkac L.M., Beanan M.J.,

RA Daugherty S.C., Deboy R.T., Durkin A.S., Kolonay J.F., Madupu R.,

RA Nelson W.C., Ayodeji B., Kraul M., Shetty J., Malek J., Van Aken

RA Riedmuller S., Tettelin H., Gill S.R., White O., Salzberg S.L.,

RA Hoover D.L., Lindler L.E., Hailing S.M., Boyle S.M., Fraser C.M.,

RT "The Brucella suis genome reveals fundamental similarities between

RT animal and plant pathogens and symbionts.";

RL Proc. Natl. Acad. Sci. U.S.A. 99:13148-13153 (2002).

DR EMBL; AE014592; AAN34160.1; -

DR TIGR; BRA0991; -

DR InterPro; IPR007916; UPF0191.

DR Pfam; PF05252; UPF0191; 1.

KW Complete proteome.

SQ SEQUENCE 220 AA; 24796 MW; AC2C060433169497 CRC64;

Query Match 2.8%; Score 8; DB 16; Length 220;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QV 54 ALLVPLAL 61

|||||

Db 132 ALLVPLAL 139

RESULT 22

Q89GW9

ID Q89GW9 PRELIMINARY; PRT; 232 AA.

AC Q89GW9;

DT 01-JUN-2003 (TrEMBLrel. 24, Created)

DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)

DE BLR6226 protein.

GN BLR6226.

OS Bradyrhizobium japonicum.

OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;

OC Bradyrhizobiaceae; Bradyrhizobium.

OX NCBI_TaxID=375;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=USDA 110;

RX MEDLINE=22484998; PubMed=12597275;

RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiumi T.,

RA Sasamoto S., Watanabe A., Idesawa K., Iriguchi M., Kawashima K.,

RA Kohara M., Matsumoto M., Shimpo S., Tsuruoka H., Wada T., Yamada

RA Tabata S.;

RT "Complete genomic sequence of nitrogen-fixing symbiotic bacteriu

RT Bradyrhizobium japonicum USDA110.";

RL DNA Res. 9:189-197 (2002).

DR EMBL; AF005957; BAC51491.1; -

DR GO; GO:0016020; C:membrane; IEA.

DR GO; GO:0005215; F:transporter activity; IEA.

DR GO; GO:0006810; P:transport; IEA.

DR InterPro; IPR000515; BPD_transp.

DR Pfam; PF00528; BPD_transp; 1.

KW Complete proteome.

SQ SEQUENCE 232 AA; 23704 MW; CD805BD1F43F1B46 CRC64;

Query Match

Best Local Similarity 100.0%; Pred. No. 32;

Matches 8; Conservative 0; Mismatches 0; Indels 0;

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RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou
RA Garber R.L., Goltry L., Tolentino E., Westbrook-Wadman S., Yuan Y
RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M
RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
RA Reizer J., Sajer M.H., Hancock R.E.W., Lory S., Olson M.V.;
RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an
RT opportunistic pathogen.";
RL Nature 406:959-964(2000).
DR EMBL; AF082575; AAC98784.1; --
DR EMBL; AF004866; AAG07909.1; --
DR FIC; D83080; D83080
DR GO; GO:0016021, C:integral to membrane; IEA.
KW Transmembrane; Complete proteome.
SQ SEQUENCE 278 AA; 30793 MW; C623F1AB0691CPEF CRC64;

Query Match 2.8%; Score 8; DB 16; Length 278;
Best Local Similarity 100.0%; Pred. No. 38;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 70 LGLLLAVV 77
Db 47 LGLLLAVV 54

RESULT 25
Q82J43 PRELIMINARY; PRT; 278 AA.
AC Q82J43;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Putative metalloproteinase.
GN SAV2939.
OS Streptomyces avermitilis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycineae; Streptomycetaceae; Streptomyces.
OC NCBI_TaxID=33903;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=21477403; PubMed=11572948;
RA Omura S., Ikeda H., Ishikawa J., Hanamoto A., Takahashi C.,
RA Shinose M., Takahashi Y., Horikawa H., Nakazawa H., Osonoe T.,
RA Kikuchi H., Shiba T., Sakaki Y., Hattori M.;
RT "Genome sequence of an industrial microorganism Streptomyces
RT avermitilis: deducing the ability of producing secondary
RT metabolites.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:12215-12220(2001).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=22608306; PubMed=12692562;
RA Ikeda H., Ishikawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba
RA Sakaki Y., Hattori M., Omura S.;
RT "Complete genome sequence and comparative analysis of the industr.
RT microorganism Streptomyces avermitilis.";
RL Nat. Biotechnol. 21:526-531(2003).
DR EMBL; AF005033; BAC70650.1; --
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0008237; F:metallopeptidase activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR000013; Peptidase_M7.
DR Pfam; PF02031; Peptidase_M7; 1.
DR PRINTS; PR00787; NEUTRALEPTASE.
DR ProDom; PD016028; Peptidase_M7; 1.
KW Complete proteome.
SQ SEQUENCE 278 AA; 28113 MW; 9545813BCAC0BFA2 CRC64;

Query Match 2.8%; Score 8; DB 16; Length 278;
Best Local Similarity 100.0%; Pred. No. 38;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 59 LALGLGLIA 66

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|||||
LGLA 20

RELIMINARY; PRT; 279 AA.

(Tremblrel. 17, Created)
(Tremblrel. 17, Last sequence update)
(Tremblrel. 23, Last annotation update)
protein.

(Mouse).
etazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
0090;

4 N.A.

/6J; TISSUE=Medulla oblongata;
3660; PubMed=11217851;
inagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
Iara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
Azaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
atsuda H.A., Ashburner M., Batalov S., Casavant T.,
J., Gaasterland T., Gissi C., King B., Kochiwa H.,
vis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
ido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
J., Bult C., Fletcher C., Carninci P., de Bonaldo M.F.,
J., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
chionni L., Mashima J., Mazzarelli I., Sakamoto N.,
Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
ato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
yo-oka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
z A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
f.;
annotation of a full-length mouse cDNA collection."
35-690(2001).
50; BAB31133.1; -
3897; 5730494G16Rik.
R002190; MAGE.
4; MAGE; 1.
0838; MAGE; 1.
79 AA; 31474 MW; 5E243590A99F15F0 CRC64;

2.8%; Score 8; DB 11; Length 279;
larity 100.0%; Pred. No. 38;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

ASLSA 87
|||||
ASLSA 48

RELIMINARY; PRT; 279 AA.

(Tremblrel. 17, Created)
(Tremblrel. 17, Last sequence update)
(Tremblrel. 23, Last annotation update)
X protein (MAGE-gi) (RIKEN CDNA 5730494G16 gene).
K.
(Mouse).
etazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
theria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
0090;

M N.A.
/6J; TISSUE=Embryo, and Embryonic stem cells;

RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.
RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush
RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Kamiya M., Lee N.
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Mazzarelli I.,
RA Lyons P., Marchionni L., Mashima J., Rodriguez I., Sakamoto N.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming
RA Wyszaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection."
RL Nature 409:685-690(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC Auquier P.H., Chomez P.M., De Backer O.R., Bertrand M.J.M.;
RT "Ten new murine members of the MAGE gene family."
RL Submitted (NOV-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=Mammary gland;
RA Strausberg R.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Head, and Spinal cord;
RL MEDLINE=22354683; PubMed=12466851;
RX The PANTOM Consortium,
RA The RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotat
RT 60,770 full-length cDNAs."
RL Nature 420:563-573(2002).
DR EMBL; AK01772; BAB30899.1; -
DR EMBL; AK010294; BAB26830.1; -
DR EMBL; AF319979; AAK01207.1; -
DR EMBL; BC034892; AAK34892.1; -
DR EMBL; AK049759; BAC33907.1; -
DR EMBL; AK076471; BAC36358.1; -
DR MGD; MGI:1913897; 5730494G16Rik.
DR InterPro; IPR002190; MAGE.
DR Pfam; PF01454; MAGE; 1.
DR PROSITE; PS50838; MAGE; 1.
SQ SEQUENCE 279 AA; 31460 MW; FE2435919BD63160 CRC64;

Query Match 2.8%; Score 8; DB 11; Length 279;
Best Local Similarity 100.0%; Pred. No. 38;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 80 GSRASLSA 87
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DB 41 GSRASLSA 48

RESULT 28

Q9VNP0
ID Q9VNP0 PRELIMINARY; PRT; 306 AA.
AC Q9VNP0; O8SZB8;
DT 01-MAY-2000 (Tremblrel. 13, Created)
DT 01-OCT-2002 (Tremblrel. 22, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE CG1169 protein (R07882p).
GN CG1169.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;

rosophilidae; Drosophila.
7;
N.A.
Y;
06; PubMed-10731132;
liniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
Lewis S.E., Li P.W., Hoskins R.A., Galie R.F.,
Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
Portman J.R., Vandeil M.D., Zhang O., Chen L.X.,
Rogers Y.H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
e C., Baxter E.G., Helt G., Nelson C.R., Miklos G.D.G.,
Bayani A., An H.-J., Andrews-Frankoch C., Baldwin D.,
asu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
enos P.V., Bertram B.P., Bhandari D., Bolshakov S.,
tchan M.R., Bouck J., Brokstein P., Brottier P.,
usam D.A., Butler H., Cadieu E., Center A., Chandra I.,
awley S., Dahike C., Davenport L.B., Davies P.,
Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
p L.E., Downes M., Dugan-Rocha S., Durkov B.C., Dunn P.,
vangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
riellian A.E., Garg N.S., Gelbart W.M., Glasser K.,
G F., Gorrell J.H., Gu Z., Guan P., Harris M.,
arvey D., Heiman T.J., Hernandez J.R., Houck J.,
ston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
ush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
odira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
B., McIntosh T.C., McLeod M.P., McPherson D.,
ilshina N.V., Mobarry C., Morris J., Moshrefi A.,
y M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
elson K.A., Nixon K., Nuskern D.R., Pacleb J.M.,
Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
ington K., Saunders R.D.C., Scheeler F., Shen H.,
en-Kiamos I., Simpson M., Skupski M.P., Smith T.,
dler A.C., Stapleton M., Strong R., Sun E.,
ector C., Turner R., Venter E., Wang A.H., Wang X.,
asarnag D.A., Weinstein G.M., Weissbach J.,
Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
ong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
ers E.W., Rubin G.M., Venter J.C.;
quence of *Drosophila melanogaster*.
85-2195(2000).
N.A.
Adams M.D., Krommiller B., Wan K.H., Holt R.A.,
ayne J.D., Amanides P.G., Brandon R.C., Rogers Y.,
H., Baldwin D., Banzon J., Beeson K.Y., Busan D.A.,
Center A., Champe M., Davenport L.B., Dietz S.M.,
sett V., Doup L.E., Doyle C., Dresnek D., Farfan D.,
rise E., Galie R.F., Garg N.S., George R.A.,
houck J., Hoskins R.A., Hostin D., Howland T.J.,
ilali M., Kruse D., Li P., Mattel B., Moshrefi A.,
Moy M., Murphy B., Nelson C., Nelson K.A., Nunoo J.,
agas V., Park S., Patel S., Pfeiffer B.,
S., Pittman G.S., Puri V., Richards S., Scheeler F.,
Strong R., Svitskas R., Tector C., Tyler D.,
Zaveri J.S., Smith H.O., Venter J.C., Rubin G.M.;
Drosophila melanogaster genome.
2-2000) to the EMBL/GenBank/DBJ databases.
N.A.
aby M.A., Matthews B.B., Bayraktaroglu L., Campbell K.,
tuang Y., Kaminker J.S., Prochuk S.E., Smith C.D.,
ngan C., Bertram B., Carlson J.W., Celniker S.E.,
sdale R., Emmert D., Frise E., de Grey A., Harris N.,
Marshall B., Millburn G., Richter J., Russo S.,
Smith E., Shu S., Smutniak F., Whitfield E.,
Gelbart W.M., Rubin G.M., Mungall C.J., Lewis S.E.;
Drosophila melanogaster genome.
2-2000) to the EMBL/GenBank/DBJ databases.
SEQUENCE FROM N.A.
Adams M.D., Celniker S.E., Gibbs R.A., Rubin G.M., Venter C.J.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
[5]
RN SEQUENCE FROM N.A.
RA FlyBase;
RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
RN [6]
RC STRAIN=Berkeley;
RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
RA Champe M., Chavez C., Dorsett V., Dresnek D., Farfan D., Frise E.,
RA George R., Gonzalez M., Guarin H., Krommiller B., Li P., Liao G.,
RA Miranda A., Mungall C.J., Nunoo J., Pacleb J., Faragas V., Park S.,
RA Patel S., Phuanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,
RA Celniker S.;
RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AS003600; AAF51889.2; -;
DR EMBL; AY070982; AAL48604.1; -;
DR FlyBase; FBgn0037428; CGI169.
SQ SEQUENCE 306 AA; 34083 MW; 32B69371475A48F9 CRC64;
Query Match 2.8%; Score 8; DB 5; Length 306;
Best Local Similarity 100.0%; Pred. No. 41;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G;
QY 56 LVPLALGL 63
Db 170 LVPLALGL 177
RESULT 29
Q864L1 PRELIMINARY; PRT; 317 AA.
ID Q864L1 AC Q864L1; DT 01-JUN-2003 (TEMBLrel. 24, Created)
DT 01-JUN-2003 (TEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TEMBLrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MC1R.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=3;
RX MEDLINE-22572539; PubMed-12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, i
RT primates."
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205086; AAP30960.1; -;
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin.
DR InterPro; IPR000276; GPCR_Rhodopsn.
DR Pfam; PF00001; 7tm.1; 1.
DR PRINTS; PR00237; GPCRHOOPS.
DR PROSITE; PS00237; G_PROTEIN_RECP_F1_1; 1.
DR PROSITE; PS00262; G_PROTEIN_RECP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34710 MW; 8815D21464BD2475 CRC64;
Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G;
QY 137 RARRATAA 144
Db 160 RARRATAA 167


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RE PRINTS; PR00237; GPCRHHODPSN.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS50262; G_PROTEIN_RECEP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34782 MW; 96A807F178FCEC21 CRC64;

Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 137 RARRAIAA 144
DB 160 RARRAIAA 167

RESULT 32
Q864J7 PRELIMINARY; PRT; 317 AA.
AC Q864J7;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MCL1.
OS Macaca silenus (Lion-tailed macaque).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheciidae;
OC Cercopitheciinae; Macaca.
OX NCBI_TaxID=54601;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1;
RX MEDLINE=22572539; PubMed=12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor,
in primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205100; AAP30974.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; P:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin.
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm_1; 1.
DR PRINTS; PR00237; GPCRHHODPSN.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS50262; G_PROTEIN_RECEP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34782 MW; 96A807F178FCEC21 CRC64;

Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 137 RARRAIAA 144
DB 160 RARRAIAA 167

RESULT 33
Q864J5 PRELIMINARY; PRT; 317 AA.
AC Q864J5;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MCL1.
OS Macaca nigra (Celebes black macaque).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheciidae;
OC Cercopitheciinae; Macaca.
OX NCBI_TaxID=54600;

```

N.A.
39; PubMed=12687585;
illy J.;
a pigmentation gene, the melanocortin-1 receptor, in
anthropol. 121:67-80(2003).
; AAP30976.1; -
; C:integral to membrane; IEA.
; F:receptor activity; IEA.
; F:rhodopsin-like receptor activity; IEA.
; P:G-protein coupled receptor protein signalin. . . ; IEA.
00276; GPCR_Rhodpsn.
7tm 1; 1.
7; GPCR_Rhodopsn.
37; G_PROTEIN_RECEP_F1_1; 1.
62; G_PROTEIN_RECEP_F1_2; 1.
AA; 34779 MW; 1A091A65BDBCBACAE CRC64;
2.8%; Score 8; DB 6; Length 317;
urity 100.0%; Pred.No. 42;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
IAAA 144
IAAA 167
ELIMINARY; PRT; 317 AA.
EMBLrel. 24, Created)
EMBLrel. 24, Last sequence update)
EMBLrel. 25, Last annotation update)
receptor.
azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Platyrrhini; Cebidae; Aotinae; Aotus.
.953;
N.A.
39; PubMed=12687585;
illy J.;
a pigmentation gene, the melanocortin-1 receptor, in
anthropol. 121:67-80(2003).
; AAP31003.1; -
; C:integral to membrane; IEA.
; F:receptor activity; IEA.
; F:rhodopsin-like receptor activity; IEA.
; P:G-protein coupled receptor protein signalin. . . ; IEA.
00276; GPCR_Rhodpsn.
7tm 1; 1.
7; GPCR_Rhodopsn.
37; G_PROTEIN_RECEP_F1_1; 1.
62; G_PROTEIN_RECEP_F1_2; 1.
AA; 34654 MW; DA5F4420DFECC4B CRC64;
2.8%; Score 8; DB 6; Length 317;
urity 100.0%; Pred.No. 42;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
IAAA 144
IAAA 167

RESULT 35
Q864G7
ID Q864G7 PRELIMINARY; PRT; 317 AA.
AC Q864G7;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MCL1.
OS Ateles paniscus (Black spider monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Ateleinae; Ate.
OX NCBI_TaxID=9510;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=3;
RX MEDLINE=22572539; PubMed=12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, in
RT primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205130; AAP31004.1; -
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin. .
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm 1; 1.
DR PRINTS; PR00237; GPCR_Rhodopsn.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS00262; G_PROTEIN_RECEP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34719 MW; 5481D6A1B9085D43 CRC64;
Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred.No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 137 RARRAIAA 144
Db 160 RARRAIAA 167
|||||
RESULT 36
Q864G6
ID Q864G6 PRELIMINARY; PRT; 317 AA.
AC Q864G6;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MCL1.
OS Alouatta seniculus (Red howler monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Alouattinae;
OC Alouatta.
OX NCBI_TaxID=9503;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=471;
RX MEDLINE=22572539; PubMed=12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, in
RT primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205131; AAP31005.1; -
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin. .
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm 1; 1.
DR PRINTS; PR00237; GPCR_Rhodopsn.

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DF 237; G_PROTEIN_RECEP_F1_1; 1.
DE 1262; G_PROTEIN_RECEP_F1_2; 1.
EQ 7 AA; 34830 MW; 87F7EFAE347671E4 CRC64;
    2.8%; Score 8; DB 6; Length 317;
    arity 100.0%; Pred. No. 42;
    conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY RAIAA 144
DE |||||
EQ RAIAA 167

RE PRELIMINARY; PRT; 317 AA.
DE (TrEMBLrel. 24, Created)
DE (TrEMBLrel. 24, Last sequence update)
DE (TrEMBLrel. 25, Last annotation update)
DE -1 receptor.
DE 1 (Bolivian red howler monkey).
DE stazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
DE Theria; Primates; Platyrrhini; Cebidae; Alouattinae;
DE ||1123;
DE 1 N.A.
DE 2539; PubMed=12687585;
DE Kelly J.;
DE a pigmentation gene, the melanocortin-1 receptor, in
DE Anthropol. 121:67-80(2003).
DE 33; AAP31007.1; -.
DE 21; C:integral to membrane; IEA.
DE 72; F:receptor activity; IEA.
DE 34; F:rhodopsin-like receptor activity; IEA.
DE 36; P:G-protein coupled receptor protein signalin...; IEA.
DE {000276; GPCR_Rhodopsn.
DE 1; 7tm 1; 1.
DE 237; GPCR_Rhodopsn.
DE 237; G_PROTEIN_RECEP_F1_1; 1.
DE 262; G_PROTEIN_RECEP_F1_2; 1.
DE 17 AA; 34686 MW; BA7B14APEC7EA971 CRC64;
    2.8%; Score 8; DB 6; Length 317;
    arity 100.0%; Pred. No. 42;
    conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY RAIAA 144
DE |||||
EQ RAIAA 167

RE PRELIMINARY; PRT; 317 AA.
DE (TrEMBLrel. 24, Created)
DE (TrEMBLrel. 24, Last sequence update)
DE (TrEMBLrel. 25, Last annotation update)
DE -1 receptor.
DE aya (Black howler monkey).
DE stazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
DE Theria; Primates; Platyrrhini; Cebidae; Alouattinae;
DE ||1123;
DE 502;

RP SEQUENCE FROM N.A.
RC STRAIN=2;
RX MEDLINE=22572539; PubMed=12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, in
RT Primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205134; AAP31009.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin.
DR InterPro; IPR000276; GPCR_Rhodopsn.
DR Pfam; PF00001; 7tm 1; 1.
DR PRINTS; PR00237; GPCR_Rhodopsn.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS00262; G_PROTEIN_RECEP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34692 MW; 91320E374CDB75DB CRC64;
    Query Match 2.8%; Score 8; DB 6; Length 317;
    Best Local Similarity 100.0%; Pred. No. 42;
    Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 137 RARRAIAA 144
DE |||||
DE 160 RARRAIAA 167

RESULT 39
Q864G2 PRELIMINARY; PRT; 317 AA.
AC Q864G2;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MC1R.
OS Alouatta palliata (Mantled howler monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Alouattinae;
OC Alouatta.
OX NCBI_TaxID=30589;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1;
RX MEDLINE=22572539; PubMed=12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, in
RT Primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205135; AAP31009.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin.
DR InterPro; IPR000276; GPCR_Rhodopsn.
DR Pfam; PF00001; 7tm 1; 1.
DR PRINTS; PR00237; GPCR_Rhodopsn.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS00262; G_PROTEIN_RECEP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34670 MW; D5414E350E2435DE CRC64;
    Query Match 2.8%; Score 8; DB 6; Length 317;
    Best Local Similarity 100.0%; Pred. No. 42;
    Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 137 RARRAIAA 144
DE |||||
DE 160 RARRAIAA 167

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ELIMINARY; PRT; 317 AA.
TREMBlrel. 24, Created)
TREMBlrel. 24, Last sequence update)
TREMBlrel. 25, Last annotation update)
receptor.

aa; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Platyrrhini; Cebidae; Alouattinae;
1253;
N.A.
339; PubMed=12687585;
illy J.;
a pigmentation gene, the melanocortin-1 receptor, in
anthropol. 121:67-80(2003).
; AAP31010.1; -.
; C: integral to membrane; IEA.
; F: receptor activity; IEA.
; F: rhodopsin-like receptor activity; IEA.
; P: G-protein coupled receptor protein signalin. . . ; IEA.
000276; GPCR_Rhodopsn.
; 7tm.1; 1.
37; GPCR_Rhodopsn.
37; G_PROTEIN_RECEP_F1_1; 1.
362; G_PROTEIN_RECEP_F1_2; 1.
; AA; 34728 MW; 976BB14FF98B4966 CRC64;
2.8%; Score 8; DB 6; Length 317;
rity 100.0%; Pred. No. 42;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
IAA 144
IAA 167
ELIMINARY; PRT; 317 AA.
TREMBlrel. 24, Created)
TREMBlrel. 24, Last sequence update)
TREMBlrel. 25, Last annotation update)
receptor.
ata rubra.
aa; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Strepsirhini; Lemuridae; Varecia.
304;
N.A.
339; PubMed=12687585;
illy J.;
a pigmentation gene, the melanocortin-1 receptor, in
anthropol. 121:67-80(2003).
; AAP31013.1; -.
; C: integral to membrane; IEA.
; F: receptor activity; IEA.
; F: rhodopsin-like receptor activity; IEA.
; P: G-protein coupled receptor protein signalin. . . ; IEA.
000276; GPCR_Rhodopsn.
; 7tm.1; 1.
37; GPCR_Rhodopsn.
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```
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
KW PROSITE; PS50262; G_PROTEIN_RECEP_F1_2; 1.
SQ RECEPTOR.
SQ SEQUENCE 317 AA; 34714 MW; C1F5DA35032717D7 CRC64;

Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 137 RARRATAA 144
Db 160 RARRATAA 167
|||||
|160 RARRATAA 167

RESULT 42
Q864F7 PRELIMINARY; PRT; 317 AA.
AC Q864F7;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MCLR.
OS Varecia variegata variegata.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Strepsirhini; Lemuridae; Varecia.
OX NCBI_TaxID=87289;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1;
RX MEDLINE=22572539; PubMed=12687585;
RA Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor, in
RT primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AX205140; AAP31014.1; -.
DR GO; GO:0016021; C: integral to membrane; IEA.
DR GO; GO:0004872; F: receptor activity; IEA.
DR GO; GO:0001584; F: rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P: G-protein coupled receptor protein signalin. .
DR InterPro; IPR000276; GPCR_Rhodopsn.
DR Pfam; PF00001; 7tm.1; 1.
DR PRINTS; PR00237; GPCR_Rhodopsn.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS50262; G_PROTEIN_RECEP_F1_2; 1.
KW RECEPTOR.
SQ SEQUENCE 317 AA; 34714 MW; C1F5DA35032717D7 CRC64;

Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 137 RARRATAA 144
Db 160 RARRATAA 167
|||||
|160 RARRATAA 167

RESULT 43
Q864F6 PRELIMINARY; PRT; 317 AA.
AC Q864F6;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Melanocortin-1 receptor.
GN MCLR.
OS Eulemur fulvus (brown lemur).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Strepsirhini; Lemuridae; Eulemur.
OX NCBI_TaxID=13515;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=2;
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539; PubMed=12687585;
Kelly J.;
a pigmentation gene, the melanocortin-1 receptor, in
Anthropol. 121:67-80(2003).
1; AAP31015.1; -.
21; C:integral to membrane; IEA.
22; F:receptor activity; IEA.
34; F:rhodopsin-like receptor activity; IEA.
36; P:G-protein coupled receptor protein signalin. . .; IEA.
0000276; GPCR_Rhodopsn.
1; 7tm1.1; -.
237; GPCR_Rhodopsn.
1237; G_PROTEIN_RECEP_F1_1; 1.
1262; G_PROTEIN_RECEP_F1_2; 1.
7 AA; 34748 MW; DAF913C3B9ECC2AF CRC64;
2.8%; Score 8; DB 6; Length 317;
arity 100.0%; Pred.No. 42;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ATAA 144
|||||
ATAA 167

PRELIMINARY; PRT; 317 AA.
(TREMBLrel. 24, Created)
(TREMBLrel. 24, Last sequence update)
(TREMBLrel. 25, Last annotation update)
-1 receptor.
seus (Gray gentle lemur) (Bamboo lemur).
stazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
heria; Primates; Strepsirhini; Lemnidae; Hapalemur.
3557;
4 N.A.
539; PubMed=12687585;
Kelly J.;
a pigmentation gene, the melanocortin-1 receptor, in
Anthropol. 121:67-80(2003).
12; AAP31016.1; -.
21; C:integral to membrane; IEA.
72; F:receptor activity; IEA.
34; F:rhodopsin-like receptor activity; IEA.
36; P:G-protein coupled receptor protein signalin. . .; IEA.
0000276; GPCR_Rhodopsn.
1; 7tm1.1; -.
237; GPCR_Rhodopsn.
1237; G_PROTEIN_RECEP_F1_1; 1.
1262; G_PROTEIN_RECEP_F1_2; 1.
17 AA; 34424 MW; 7FB912CCC3F5EC71 CRC64;
2.8%; Score 8; DB 6; Length 317;
arity 100.0%; Pred.No. 42;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ATAA 144
|||||
ATAA 167

PRELIMINARY; PRT; 317 AA.

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Q864F4;
01-JUN-2003 (TREMBLrel. 24, Created)
01-JUN-2003 (TREMBLrel. 24, Last sequence update)
01-OCT-2003 (TREMBLrel. 25, Last annotation update)
Melanocortin-1 receptor.
MCLR.
Lemur catta (Ring-tailed lemur).
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
OC Mammalia; Eutheria; Primates; Strepsirhini; Lemnidae; Lemur.
NCBI_TaxID=9447;
(1)
RN SEQUENCE FROM N.A.
RP STRAIN=3;
RC MEDLINE=22572539; PubMed=12687585;
RX Mundy N.I., Kelly J.;
RT "Evolution of a pigmentation gene, the melanocortin-1 receptor,
RT primates.";
RL Am. J. Phys. Anthropol. 121:67-80(2003).
DR EMBL; AY205143; AAP31017.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0004872; F:receptor activity; IEA.
DR GO; GO:0001584; F:rhodopsin-like receptor activity; IEA.
DR GO; GO:0007186; P:G-protein coupled receptor protein signalin.
DR InterPro; IPR000276; GPCR_Rhodopsn.
DR Pfam; PF00001; 7tm1.1; -.
DR PRINTS; PR00237; GPCR_Rhodopsn.
DR PROSITE; PS00237; G_PROTEIN_RECEP_F1_1; 1.
DR PROSITE; PS0262; G_PROTEIN_RECEP_F1_2; 1.
KW Receptor.
SQ SEQUENCE 317 AA; 34667 MW; 3E7419FDEC2DE738 CRC64;

Query Match 2.8%; Score 8; DB 6; Length 317;
Best Local Similarity 100.0%; Pred.No. 42;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 137 RARRAIAA 144
DB 160 RARRAIAA 167
|||||

RESULT 46
OS8554 PRELIMINARY; PRT; 339 AA.
ID OS8554;
AC OS8554;
DT 01-AUG-1998 (TREMBLrel. 07, Created)
DT 01-JAN-1999 (TREMBLrel. 09, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
DE Hypothetical protein PH0824.
GN PH0824.
OS Pyrococcus horikoshii.
OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococca
OC Pyrococcus.
OX NCBI_TaxID=53953;
(1)
RN SEQUENCE FROM N.A.
RP STRAIN=OT3;
RC MEDLINE=98344137; PubMed=9679194;
RA Kawarabayashi Y., Sawada M., Horikawa H., Haikawa Y., Hino Y.,
RA Yamamoto S., Sekine M., Baba S.-I., Kosugi H., Hosoyama A., Naga
RA Sakai M., Ogura K., Otsuka R., Nakazawa H., Takamiya M., Ohfuku
RA Fundashii T., Tanaka T., Kudoh Y., Yamazaki J., Kushida N., Oguci
RA Aoki K.-I., Yoshizawa T., Kuchimura Y., Robb F.T., Horikoshi K.,
RA Masuchi Y., Shizuya H., Kikuchi H.;
RT "Complete sequence and gene organization of the genome of a hype-
RL DNA Res. 5:55-76(1998).
DR EMBL; AP000003; BAA29917.1; -.
DR FIR; C71132; C71132.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0008508; F:bile acid:sodium symporter activity; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR002657; BilAC/Na_symport.
DR Pfam; PF01758; SBF; 1.

```

rotein; Complete proteome.
AA; 37228 MW; E91697D5C8C3705F CRC64;
2.8%; Score 8; DB 17; Length 339;
rity 100.0%; Pred. No. 45;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
AWV 77
|||
AVV 118
ELIMINARY; PRT; 342 AA.
REMBLrel. 03, Created)
REMBLrel. 03, Last sequence update)
REMBLrel. 24, Last annotation update)
Human).
azo; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Catarrhini; Hominidae; Homo.
6;
N.A.
28; PubMed-8702217;
on N., Lee S.W., Liebert M., Grossman H.B.;
lysis of a gene, Bb1, overexpressed in bladder and
ma.";
16:1085-1090 (1996).
AAB37433.1; -;
04299; MBOAT_fam.
MBOAT; 1.
AA; 38163 MW; 2B479EA8CF1B91C CRC64;
2.8%; Score 8; DB 4; Length 342;
rity 100.0%; Pred. No. 45;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
LAL 67
|||
LAL 317
ELIMINARY; PRT; 343 AA.
REMBLrel. 17, Created)
REMBLrel. 17, Last sequence update)
REMBLrel. 22, Last annotation update)
rotein.
Human).
azo; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Catarrhini; Hominidae; Homo.
6;
N.A.
and Colon;
3-2001) to the EMBL/GenBank/DBJ databases.
; AAH03164.1; -;
; AAH02512.1; -;
04299; MBOAT_fam.
MBOAT; 1.
rotein.
AA; 38727 MW; F71E7DBF74BD9BB7 CRC64;
2.8%; Score 8; DB 4; Length 343;
rity 100.0%; Pred. No. 45;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 ALGLGLAL 67
|||
Db 311 ALGLGLAL 318
|||
RESULT 49
Q8TUU8
ID Q8TUU8 PRELIMINARY; PRT; 370 AA.
AC Q8TUU8;
DT 01-JUN-2002 (TREMBLrel. 21, Created)
DT 01-JUN-2002 (TREMBLrel. 21, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
DE Permease subunit of a ABC-type transport system involved in
DE lipoprotein release.
GN MK1655
OS Methanopyrus kandleri.
OC Archaea; Euryarchaeota; Methanopyri; Methanopyrales; Methanopyrac
OC Methanopyrus.
OX NCBI_TaxID=2320;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=AV19 / DSM 6324 / JCM 9639;
RX MEDLINE=21927647; PubMed=11930014;
RA Slesarev A.I., Mezheva V.V., Makarova K.S., Polushin N.N.,
RA Shcherbinina O.V., Shakhova V.V., Belova G.I., Aravind L.,
RA Natale D.A., Rogozin I.B., Tatusov R.L., Wolf Y.I., Stetter K.O.,
RA Malykh A.G., Koonin E.V., Kozlyavkin S.A.;
RT "The complete genome of hyperthermophile Methanopyrus kandleri AV.
RT and monophyly of archaeal methanogens.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:4644-4649 (2002).
DR EMBL; AB010455; AAM02868.1; -;
DR GO; GO:0016020; C:membrane; IEA.
DR InterPro; IPR003838; DUF214.
DR Pfam; PF02687; FtsX; 1.
KW Complete proteome.
SQ SEQUENCE 370 AA; 39411 MW; B07662EALB5A644E CRC64;
Query Match 2.8%; Score 8; DB 17; Length 370;
Best Local Similarity 100.0%; Pred. No. 49;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G
QY 61 LGLGLALA 68
|||
Db 336 LGLGLALA 343
|||
RESULT 50
O53860
ID O53860 PRELIMINARY; PRT; 372 AA.
AC O53860;
DT 01-JUN-1998 (TREMBLrel. 06, Created)
DT 01-JUN-1998 (TREMBLrel. 06, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
DE Hypothetical protein cysM3.
GN CYSM3 OR RV0848 OR MTV043.41.
OS Mycobacterium tuberculosis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37Rv;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris
Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekai F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd
Hornsbay T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrett B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the

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me sequence." ;
7-544(1998) ;
4; CAA17654.1; -.
H70813.
1JBQ.
RV0848; -.
9; P-lyase activity; IEA.
0; P-lyase activity; IEA.
001926; B6_enzyme_beta.
; PALP; 1.
Protein; Complete proteome.
2 AA; 40118 MW; 927386BEDI5FB6C CRC64;
2.8%; Score 8; DB 16; Length 372;
arity 100.0%; Pred. No. 49;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
HALA 68
|||||
HALA 103

PRELIMINARY; PRT; 372 AA.

TREMBLrel. 25, Created)
TREMBLrel. 25, Last sequence update)
TREMBLrel. 25, Last annotation update)
eine synthase A CYSK2 (O-acetylserine sulphydrylase)
ne (Thiol)-lyase) (CSASE) (EC 2.5.1.47).
71.
bovis.
inobacteria; Actinobacteridae; Actinomycetales;
neae; Mycobacteriaceae; Mycobacterium.
65;
[N.A.
/97;
107; PubMed=12788972;
igleier K., Camus J.-C., Medina N., Mansoor H.,
boy S., Grondin S., Lacroix C., Monsemp C., Simon S.,
Kin R., Doggett J., Mayes R., Keating L., Wheeler P.R.,
Barrell B.G., Cole S.T., Gordon S.V., Hewinson R.G.;
genome sequence of Mycobacterium bovis." ;
acad. Sci. U.S.A. 100:7877-7882(2003).
16; CAD93733.1; -.
Complete proteome.
12 AA; 40106 MW; 6PD459DFA6FB4284 CRC64;
2.8%; Score 8; DB 16; Length 372;
arity 100.0%; Pred. No. 49;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
HALA 68
|||||
HALA 103

PRELIMINARY; PRT; 375 AA.

(TREMBLrel. 24, Created)
(TREMBLrel. 24, Last sequence update)
(TREMBLrel. 25, Last annotation update)
irogenase beta subunit.
3 maripaludis.
archaeota; Methanococci; Methanococcales;
seae; Methanococcus.
9152;

SEQUENCE FROM N.A.
STRAIN=LL/S2;
RC MEDLINE=22557897; PubMed=12670979;
RA Wood G.E., Haydock A.K., Leigh J.A.;
RT "Function and Regulation of the Formate Dehydrogenase Genes of th
RI Methanogenic Archaeon Methanococcus maripaludis." ;
RL J. Bacteriol. 185:2548-2554(2003).
DR EMBL; AY236516; AA085931.1; -.
DR GO; GO:0005489; P:electron transport; IEA.
DR GO; GO:0006118; P:electron transport; IEA.
DR InterPro; IPR001450; 4Pe4S_ferredoxin.
DR InterPro; IPR007525; Fhb_FdhB_C.
DR InterPro; IPR007516; Fhb_FdhB_N.
DR Pfam; PF00037; feir4; 1.
DR Pfam; PF04432; Fhb_FdhB_C; 1.
DR Pfam; PF04422; Fhb_FdhB_N; 1.
DR PROSITE; PS00198; 4Fe4S_FERREDOXIN; 2.
SQ SEQUENCE 375 AA; 42431 MW; AB09678918AEC8AD CRC64;

Query Match 2.8%; Score 8; DB 16; Length 375;
Best Local Similarity 100.0%; Pred. No. 49;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 217 LVDGVIAL 224
DB 35 LVDGVIAL 42

RESULT 53
Q9KVT3 PRELIMINARY; PRT; 377 AA.
ID Q9KVT3
AC Q9KVT3;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Putative integral membrane protein.
GN SC05682 OR SC5H4.06C.
OS Streptomyces coelicolor.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycineae; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=1902;
RN [1]
SEQUENCE FROM N.A.
RC STRAIN=A3(2) / M145;
RX MEDLINE=21996410; PubMed=12000953;
RA Bentley S.D., Chater K.P., Cerdeno-Tarraga A.-M., Challis G.L.,
RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,
RA Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,
RA Cronin A., Fraser A., Goble A., Hidalgo J., Hornsby T., Howarth S.,
RA Huang C.-H., Kieser T., Larke L., Murphy L., Oliver K., O'Neill S.,
RA Rabinowitsch E., Rajandream M.A., Rutherford K., Rutter S.,
RA Seeger K., Saunders D., Sharp S., Squares R., Squares S., Taylor
RA Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,
RA Hopwood D.A.;
RT "Complete genome sequence of the model actinomycete Streptomyces
coelicolor A3(2).";
RL Nature 417:141-147(2002).
DR EMBL; AL939124; CAB91118.1; -.
KW Complete proteome.
SQ SEQUENCE 377 AA; 37614 MW; A35DA0437F044C45 CRC64;

Query Match 2.8%; Score 8; DB 16; Length 377;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 59 LALGUGLA 66
DB 327 LALGUGLA 334

RESULT 54
Q9SHD8 PRELIMINARY; PRT; 387 AA.
ID Q9SHD8

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REMBLrel. 13, Created)
 REMBLrel. 13, Last sequence update)
 REMBLrel. 25, Last annotation update)
 1.20 (At2g45000 protein).
 T2G45000.
 aliana (Mouse-ear cress).
 idiplantae; Streptophyta; Tracheophyta;
 Magnoliophyta; eudicotyledons; core eudicots; rosids;
 rassicales; Brassicaceae; Arabidopsis.
 2;
 N.A.
 H., Cheuk R., Kim C.J., Lim J., Meyers M.C., Banh J.,
 ninci P., Chang E., Dale J.M., Goldsmith A.D.,
 Ishida J., Jones T., Kamiya A., Karlin-Neumann G.,
 B., Lee J.M., Lin J., Miranda M., Narusaka M.,
 dera C.S., Palm C.J., Quach H.L., Sakurai T., Satou M.,
 wick A., Tang C.C., Toriumi M., Wu H.C., Yamada K.,
 u G., Yu S., Shinozaki K., Davis R.W., Theologis A.,
 DNA clones.";
 -2002) to the EMBL/GenBank/DBJ databases.
 N.A.
 S.X., Sakano H., Pham P.K., Banh J., Chung M.K.,
 Lee J.M., Quach H.L., Toriumi M., Yu G., Bowser L.,
 hen H., Cheuk R., Hayashizaki Y., Ishida J., Jones T.,
 lin-Neumann G., Kawai J., Kim C., Lam B., Lin J.,
 rusa M., Nguyen M., Palm C.J., Sakurai T., Satou M.,
 P., Southwick A., Shinozaki K., Davis R.W., Ecker J.R.,
 ull Length cDNA Clones";
 -2002) to the EMBL/GenBank/DBJ databases.
 N.A.
 umbia;
 87; PubMed=10617197;
 , Rounsley S.D., Shea T.P., Benito M.-I., Town C.D.,
 son T.W., Bowman C.L., Barnstead M.E., Feidblyum T.V.,
 tchum K.A., Lee J.J., Rinning C.M., Koo H., Moffat K.S.,
 hen M., Vanaken S.E., Unayam L., Tallon L.J., Gill J.R.,
 riera A.J., Creasy T.H., Goodman H.M., Somerville C.R.,
 , Preuss D., Nierman W.C., White O., Eisen J.A.,
 Fraser C.M., Venter J.C.;
 analysis of chromosome 2 of the plant Arabidopsis
 -768(1999).
 N.A.
 umbia;
 -2000) to the EMBL/GenBank/DBJ databases.
 ; AAL69462.1; -;
 ; AAL86303.1; -;
 ; AAD32835.1; -;
 84885.
 xotein.
 AA; 40584 MW; AF6C6B3BAC9BF69A CRC64;
 2.8%; Score 8; DB 10; Length 387;
 rity 100.0%; Pred. No. 51;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 EED 100
 ||||
 .EED 377
 ELIMINARY; PRT; 397 AA.
 85
 86
 87
 88
 89
 90
 91
 92
 93
 94
 95
 96
 97
 98
 99
 100

DT 01-JUN-2002 (TREMELrel. 21, Created)
 DT 01-JUN-2002 (TREMELrel. 21, Last sequence update)
 DT 01-JUN-2003 (TREMELrel. 24, Last annotation update)
 DE Hypothetical protein Atu3948.
 GN ATU3948 OR AGR_L_1808.
 OS Agrobacterium tumefaciens (strain C58 / ATCC 33970).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
 OC Rhizobiaceae; Rhizobium/Agrobacterium group; Agrobacterium.
 OX NCBI_TaxID=176299;
 RN [1]_TaxID=176299;
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21608550; PubMed=11743193;
 RA Wood D.W., Setubal J.C., Kaul R., Monks D.E., Kitajima J.P.,
 Okura V.K., Zhou Y., Chen L., Wood G.E., Almeida N.F. Jr., Woo L.,
 Chen Y., Paulsen I.T., Eisen J.A., Karp P.D., Bovee D. Sr.,
 Chapman P., Clendenning J., Deatherage G., Gillet W., Grant C.,
 Kutyavina T., Levy R., Li M.-J., McClelland E., Palmeri A., Gordon
 Raymond C., Rouse G., Saenphimmachak C., Wu Z., Romero P., Perry M.,
 Zhang S., Yoo H., Tao Y., Biddle P., Jung M., Krespan W., Perry M.,
 Gordon-Kamm B., Liao L., Kim S., Hendrick C., Zhao Z.-Y., Dolan M.,
 Chumley F., Tingey S.V., Tomb J.-F., Gordon M.P., Olson M.V.,
 RA Nester E.W.;
 RT "The genome of the natural genetic engineer Agrobacterium tumefaci
 RT C58.";
 RL Science 294:2317-2323(2001).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21608551; PubMed=11743194;
 RA Goodner B., Hinkle G., Gattung S., Miller N., Blanchard M., Mullin
 Qurrello B., Goldman B.S., Cao Y., Askenazi M., Halling C., Mullin
 RA Hummel K., Gordon J., Vaudin M., Iartchouk O., Epp A., Liu F.,
 RA Wollam C., Allinger M., Doughty D., Scott C., Lappas C., Markelz E
 RA Planagan C., Crowell C., Gurson J., Lomo C., Sear C., Strub G.,
 Cielo C., Slater S;
 RT "Genome sequence of the plant pathogen and biotechnology agent
 RT Agrobacterium tumefaciens C58.";
 RL Science 294:2323-2328(2001).
 DR EMBL; AE009325; AAL44750.1; ALT_INIT.
 DR EMBL; AE008289; AAK89478.1; -;
 DR PIR; AH3041; AH3041.
 DR PIR; D98244; D98244.
 DR InterPro; IPR001220; Lectin legB.
 DR InterPro; IPR001608; UPF0001.
 DR Pfam; PF01168; Ala racemase N; 1.
 DR PROSITE; PS00307; LECTIN LEGUME BETA; 1.
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 397 AA; 41708 MW; 700748E32A46AE86 CRC64;
 Query Match 2.8%; Score 8; DB 16; Length 397;
 Best Local Similarity 100.0%; Pred. No. 52;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps
 QY 251 LALRPGSS 258
 DB. |||||
 341 LALRPGSS 348
 RESULT 56
 QYU9H4
 ID QYU9H4 PRELIMINARY; PRT; 431 AA.
 AC QYU9H4;
 DT 01-OCT-2003 (TREMELrel. 25, Created)
 DT 01-OCT-2003 (TREMELrel. 25, Last sequence update)
 DT 01-OCT-2003 (TREMELrel. 25, Last annotation update)
 DE Possible bicarbonate transporter, ICT family.
 GN SYNW0284.
 OS Synechococcus sp. (strain WH8102).
 OC Bacteria; Cyanobacteria; Chroococcales; Synechococcus.
 OX NCBI_TaxID=84588;
 RN [1]_TaxID=84588;
 RP SEQUENCE FROM N.A.
 RX MEDLINE=22825697; PubMed=12917641;
 RA Palenik B., Brahamsha B., Larimer F.W., Land M., Hauser L., Chain

Regala W., Allen E.E., McCarren J., Paulsen I.,
Partensky F., Webb E.A., Waterbury J.;
if a motile marine Synechococcus";
37-1042(2003).
9; CA506799.1; -.
eome.
1 1 AA; 46300 MW; 66295F913903DBAE CRC64;
arity 2.8%; Score 8; DB 16; Length 431;
arity 100.0%; Pred. No. 56;
conservative 0; Mismatches 0; Indels 0; Gays

RELIMINARY; PRT; 435 AA.

TREMBLrel. 20, Created)
TREMBLrel. 20, Last sequence update)
TREMBLrel. 25, Last annotation update)
smbrane protein.
.03756.
anacearum (Pseudomonas solanacearum).
lasmid.
teobacteria; Betaproteobacteria; Burkholderiales;
eae; Ralstonia.
5;
{ N.A.
10;
879; PubMed:11823852;
; Genin S., Artiguenave F., Gouzy J., Mangenot S.,
lault A., Brottier P., Camus J.C., Cattolico L.,
Choisne N., Claudel-Renard C., Cunnc S., Demange N.,
vie M., Moisan A., Robert C., Saurin W., Schiex T.,
heault P., Whalen M., Wincker P., Levy M.,
nce of the plant pathogen Ralstonia solanacearum."
7-502(2002).
9; CAD17762.1; -.
1; C:extrachromosomal DNA; IEA.
4; F:ATP binding; IEA.
0; F:nucleoside-diphosphate kinase activity; IEA.
1; P:CTP biosynthesis; IEA.
3; P:GTP biosynthesis; IEA.
8; P:UTP biosynthesis; IEA.
001564; NDK.
469; NDP_KINASRS; 1.
5 AA; 47048 MW; CCB859D9C54DDB5A CRC64;
lete proteome.

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2.8%; Score 8; DB 16; Length 435;
arity 100.0%; Pred. NO. 56;
conservative 0; Mismatches 0; Indels 0; Gaps 0;

IVLAL 224
|||||
IVLAL 365

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PRELIMINARY;          PRT;   465 AA.

..TremBLrel. 20, Created)
..TremBLrel. 20, Last sequence update)
..TremBLrel. 25, Last annotation update)
..Tran protein (Putative permease, major facilitator

```

CN	YEGB OR YFO2850 OR Y1393.	
OS	Yersinia pestis.	
OC	Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;	
OC	Enterobacteriaceae; Yersinia.	
NCBI	TaxID=632;	
[1]_	NCBI_TaxID=632;	
RP	SEQUENCE FROM N.A.	
RP	STRAIN-CO-92 / Biovar Orientalis;	
RC	MEDLINE=21470413; PubMed=11586360;	
RX	Parhill J., Wren B.W., Thomson N.R., Titball R.W., Holden M.T.G.	
RA	Prentice M.B., Sebailia M., James K.D., Churcher C., Mungall K.L.	
RA	Baker S., Basham D., Bentley S.D., Brooks K., Cerdeno-Tarraga A.N.	
RA	Chillingworth T., Cronin A., Davies R.M., Davis P., Dougan G.	
RA	Felwell T., Hanlin N., Holroyd S., Jagels K., Karlyshev A.V.	
RA	Leather S., Moule S., Oyston P.C.F., Quail M., Rutherford K.	
RA	Simmonds M., Skelton J., Stevens K., Whitehead S., Barrall B.G.;	
RT	"Genome sequence of Yersinia pestis, the causative agent of plague	
RL	Nature 413:523-527(2001).	
[2]		
RP	SEQUENCE FROM N.A.	
RP	STRAIN-KIMS / Biovar Mediaevalis;	
RC	MEDLINE=22137863; PubMed=12142430;	
RX	Deng W., Burland V., Plunkett G. III, Boutin A., Mayhew G.F., Liu	
RA	Perna N.T., Rose D.J., Mau B., Zhou S., Schwartz D.C.,	
RA	Fetherston J.D., Lindler L.E., Brubaker R.R., Plano G.V.,	
RA	Straley S.C., McDonough K.A., Nilles M.L., Matson J.S., Blattner	
RA	Perry R.D.;	
RT	"Genome sequence of Yersinia pestis KIM. ";	
RL	J. Bacteriol. 184:4601-4611(2002).	
DR	EMBL; A0414154; CAC92102.1; -	
DR	EMBL; AF013741; AAM84955.1; -	
DR	PIR; AC0347; AC0347.	
DR	GO; GO:0016031; C:integral to membrane; IEA.	
DR	GO; GO:0015520; P:tetracycline:hydrogen antiporter activity; IEA	
DR	GO; GO:0005215; P:transporter activity; IEA.	
DR	GO; GO:0035904; P:tetracycline transport; IEA.	
DR	GO; GO:0006810; P:transport; IEA.	
DR	InterPro; IPR007114; MFS.	
DR	InterPro; IPR005828; Sub transporter.	
DR	InterPro; IPR001411; TCR_TetB.	
DR	Pfam; PF00083; sugar tr_1.	
DR	PRINTS; PR01036; TCTETB.	
DR	PROSITE; PS00850; MFS; 1.	
KW	Hypothetical protein; Complete proteome.	
QY	SEQUENCE 465 AA; 50176 MW; 0CC273F10B83F5ED CRC64;	
QY	Query Match 2.8%; Score 8; DB 16; Length 465;	
Db	Best Local Similarity 100.0%; Pred. No. 60;	
Db	Matches 8; Conservative 0; Mismatches 0; Indels 0;	
QY	72 LLLAVVSL 79	
Db	335 LLLAVVSL 342	
RESULT 59		
Q9RR18	PRELIMINARY; PRT; 471 AA.	
ID	Q9RR18	
AC	Q9RR18;	
DT	01-MAY-2000 (TREMBlrel. 13, Created)	
DT	01-MAY-2000 (TREMBlrel. 13, Last sequence update)	
DE	01-JUN-2003 (TREMBlrel. 24, Last annotation update)	
DE	Transport protein, putative.	
GN	DR2502.	
OC	Deinococcus radiodurans.	
OC	Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;	
OC	Deinococcaceae; Deinococcus.	
NCBI	TaxID=1299;	
[1]_	NCBI_TaxID=1299;	
RP	SEQUENCE FROM N.A.	
RP	STRAIN-R1 / ATCC 13939 / DSM 20539 / NCIB 9279;	
RX	MEDLINE=20036896; PubMed=10567266;	
RA	White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.	

aft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
in H., Jiang L., Pamphile W., Crosby M., Shen M.,
Lam P., McDonald L., Utterback T., Zalewski C.,
Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
et al. The genome sequence of the radioreistant bacterium Deinococcus
"J";
71-1577(1999).
; AAF12043.1; -.
75267.
-.

ome.
AA; 47974 MW; 96B2BBBF6E445D27 CRC64;

2.8%; Score 8; DB 16; Length 471;

ity 100.0%; Pred. No. 61;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

HAL 67

|||||

HAL 373

ELIMINARY; PRT; 472 AA.

EMBLrel. 19, Created)

EMBLrel. 19, Last sequence update)

EMBLrel. 22, Last annotation update)

rotein FLJ31346.

Human).

azoa; Chordata; Craniata; Vertebrata; Euteleostomi;
eria; Primates; Catarrhini; Hominidae; Homo.
6;

N.A.

mazaki M., Watanabe K., Kumagai A., Itakura S.,

ujimori Y., Komiyama M., Sugiyama T., Irie R.,

o H., Wakatsuki S., Ishii S., Yamamoto J., Isono Y.,

Saito K., Nishikawa T., Kimura K., Yamashita H.,

amuro Y., Sekine M., Kikuchi H., Kanda K., Wagatsuma M.,

anehori K., Takahashi-Fujii A., Oshima A., Sugiyama A.,

uzuki Y., Sugano S., Nagahari K., Masuho Y., Nagai K.,

et al. DNA sequencing project."

2001) to the EMBL/GenBank/DBJ databases.

; BAB71043.1; -.

04299; MBOAT_fam.

protein; 1.

; AA; 52774 MW; EA721998043F9EBD CRC64;

HAL 67

|||||

HAL 447

2.8%; Score 8; DB 4; Length 472;

ity 100.0%; Pred. No. 61;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

ELIMINARY; PRT; 473 AA.

EMBLrel. 21, Created)

EMBLrel. 21, Last sequence update)

EMBLrel. 22, Last annotation update)

30589L02 gene.

(Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Strausberg R.;
RL Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC023417; AAH23417.1; -.
DR MGD; MGI:1924832; 5730589L02Rik.
DR InterPro; IPR004299; MBOAT_fam.
DR Pfam; PF03062; MBOAT; 1
SQ SEQUENCE 473 AA; 53382 MW; DAALFEODA78013EA CRC64;

Query Match 2.8%; Score 8; DB 11; Length 473;

Best Local Similarity 100.0%; Pred. No. 61;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps

QY 65 LALACLG 72

|||||

Db 436 LALACLG 443

RESULT 62

Q9CY76

AC Q9CY76 PRELIMINARY; PRT; 473 AA.

DT 01-JUN-2001 (TrEMBLrel. 17, Created)

DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)

DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)

DE 5730589L02Rik protein.

GN 5730589L02Rik.

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus

OX NCBI_TaxID=10090;

RN [1]

RP SEQUENCE FROM N.A.

STRAIN=C57BL/6J; TISSUE=Embryo;

RC MEDLINE=21085660; PubMed=11217851;

RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.

RA Arakawa T., Hara A., Fukunishi Y., Kono H., Adachi J., Fukuda S.

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.

RA Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito R.

RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,

RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,

RA Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush

RA Schraml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio

RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,

RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.P.,

RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,

RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H

RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,

RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,

RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.

RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilmit

RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawai H., Kohtsuki S.

RA Hayashizaki Y.

RT "Functional annotation of a full-length mouse cDNA collection.";

RL Nature 409:685-690(2001).

DR EMBL; AK019981; BAB31950.1; -.

DR MGD; MGI:1924832; 5730589L02Rik.

DR InterPro; IPR004299; MBOAT_fam.

DR Pfam; PF03062; MBOAT; 1.

SQ SEQUENCE 473 AA; 53504 MW; CE6F8E93C3D01C4F CRC64;

Query Match 2.8%; Score 8; DB 11; Length 473;

Best Local Similarity 100.0%; Pred. No. 61;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps

QY 65 LALACLG 72

|||||

Db 436 LALACLG 443

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RE Q9LIW0 PRELIMINARY; PRT; 473 AA.
AC Q9LIW0;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
DE Similar to an Arabidopsis thaliana chromosome BAC genomic
DE sequence.
OS Oryza sativa (Rice).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=4530;
RN [1]
RP SEQUENCE FROM N.A.
RA Hsing Y.C., Chow T., Chen C., Wu H., Chu M., Chao Y., Liu S.;
RT "Oryza sativa PAC P6699E04 genomics sequence, complete sequence."
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP001111; BAA90509.1; -.
DR Gramene; Q9LIW0; -.
DR SEQUENCE 522 AA; 54697 MW; 21C6BAD2441B56BF CRC64;
SQ SEQUENCE 522 AA; 54697 MW; 21C6BAD2441B56BF CRC64;

Query Match 2.8%; Score 8; DB 10; Length 522;
Best Local Similarity 100.0%; Pred. No. 67;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 43 RRRGRGGE 50
DB 415 RRRGRGGE 422

RESULT 65
Q9LIW0 PRELIMINARY; PRT; 522 AA.
AC Q9LIW0;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
DE Similar to an Arabidopsis thaliana chromosome BAC genomic
DE sequence.
OS Oryza sativa (Rice).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=4530;
RN [1]
RP SEQUENCE FROM N.A.
RA Hsing Y.C., Chow T., Chen C., Wu H., Chu M., Chao Y., Liu S.;
RT "Oryza sativa PAC P6699E04 genomics sequence, complete sequence."
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP001111; BAA90509.1; -.
DR Gramene; Q9LIW0; -.
DR SEQUENCE 522 AA; 54697 MW; 21C6BAD2441B56BF CRC64;
SQ SEQUENCE 522 AA; 54697 MW; 21C6BAD2441B56BF CRC64;

Query Match 2.8%; Score 8; DB 10; Length 522;
Best Local Similarity 100.0%; Pred. No. 67;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 43 RRRGRGGE 50
DB 415 RRRGRGGE 422

RESULT 66
Q7WZ71 PRELIMINARY; PRT; 535 AA.
AC Q7WZ71;
DT 01-OCT-2003 (TRENBLrel. 25, Created)
DT 01-OCT-2003 (TRENBLrel. 25, Last sequence update)
DE Putative mannosyltransferase.
GN DBV20.
OS Nonomuraea sp. ATCC 39727.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptosporangineae; Streptosporangiaceae; Nonomuraea.
OX NCBI_TaxID=93944;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-ATCC 39727;
RA Sosio M., Stinchl S., Beltrametti F., Lazzarini A., Donadio S.;
RT "The gene cluster for the biosynthesis of the glycopeptide antibi-
RT A40926 by Nonomuraea sp.";
RL Submitted (MAY-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ561198; CAD91215.1; -.
KW Acyltransferase; Glycosyltransferase; Monooxygenase; Transferase
SQ SEQUENCE 535 AA; 57418 MW; 3C9059338B3308AC CRC64;

Query Match 2.8%; Score 8; DB 2; Length 535;
Best Local Similarity 100.0%; Pred. No. 68;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 122 RLVRPRRS 129
DB 480 RLVRPRRS 487

RESULT 67
Q8PMH8 PRELIMINARY; PRT; 537 AA.
AC Q8PMH8;
DT 01-OCT-2002 (TRENBLrel. 22, Created)
DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)
DT 01-JUN-2003 (TRENBLrel. 24, Last annotation update)
DE Oligopeptide transporter.

```

0. onopodis (pv. citri).
 obacteria; Gammaproteobacteria; Xanthomonadales;
 ae; Xanthomonas.
 29;
 N.A.
 TCC 13902 / XV 101;
 45; PubMed12024217;
 ,; Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,
 Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
 do Amaral A.M., Bertolini M.C., Camargo L.B.A.,
 Cannavan F., Cardoso J., Chambergo F., Chapina L.P.,
 B., Coutinho L.B., Cursino-Santos J.R., El-Dorri H.,
 rreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
 F., Franco M.C., Greggio C.C., Gruber A.,
 Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
 Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
 Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
 Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
 Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
 Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
 Santos M., Truffi D., Tsai S.M., White F.F.,
 Kitajima J.P.;
 the genomes of two Xanthomonas pathogens with differing
 ties.";
 1-463 (2002).
 ; AAM36320.1; -;
 ; C-membrane; IEA.
 ; F-transporter activity; IEA.
 ; Polipeptide transport; IEA.
 100109; PTR2.
 PTR2; 1.
 some.
 ; AA; 58369 MW; 7844COCOFEE8670 CRC64;
 2.8%; Score 8; DB 16; Length 537;
 rity 100.0%; Pred. No. 68;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 VVS 78
 ||||
 VVS 263
 ELIMINARY; PRT; 564 AA.
 REMBLrel. 10, Created)
 REMBLrel. 10, Last sequence update)
 REMBLrel. 25, Last annotation update)
 H5 (Fragment).
 A.
 negative-strand viruses; Orthomyxoviridae;
 ruses.
 911;
 N.A.
 Potsdam/2216-4/84;
 02; PubMed9882316;
 ; Zhou N., Kawoka Y., Webster R.;
 glycoproteins of H5 influenza viruses isolated from
 ns, and wild aquatic birds have distinguishable
 146-1155(1999).
 HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 TORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 OMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (A2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 ; BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 ; AAD13573.1; -;
 1HTM.

DR GO; GO:0019031; C:viral envelope; IEA.
 DR InterPro; IPR008980; Capsid hemag.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR ProDom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON TER 564 564
 SQ SEQUENCE 564 AA; 63562 MW; B317179A7F3E6F98 CRC64;
 Query Match 2.8%; Score 8; DB 12; Length 564;
 Best Local Similarity 100.0%; Pred. No. 71;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G
 QY 72 LLLAVVSL 79
 |||||
 Db 6 LLLAVVSL 13
 RESULT 69
 Q8JN92 PRELIMINARY; PRT; 568 AA.
 AC Q8JN92;
 DT 01-OCT-2002 (TrEMBLrel. 22, Created)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Hemagglutinin H5.
 GN H5.
 OS Influenza A virus (A/Goose/Hong Kong/3014.5/2000(H5N1)).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCBI_TaxID=186167;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Goose/Hong Kong/3014.5/2000;
 RX MEDLINE=22016166; PubMed=12021367;
 RA Tumpey T.M., Suarez D.L., Perkins L.E.L., Senne D.A., Lee J.G.,
 Lee Y.J., Mo I.P., Sung H.W., Swayne D.E.;
 RT "Characterization of a Highly Pathogenic H5N1 Avian Influenza A V:
 RT Isolated from Duck Meat";
 RL J. Virol. 76:6344-6355(2002).
 CC -!- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRU:
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -!- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CH:
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -!- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AY075030; AAL75843.1; -;
 DR GO; GO:0019031; C:viral envelope; IEA.
 DR InterPro; IPR008980; Capsid hemag.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR ProDom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 568 AA; 64244 MW; E0D741A75C6E76FC CRC64;
 Query Match 2.8%; Score 8; DB 12; Length 568;
 Best Local Similarity 100.0%; Pred. No. 72;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G
 QY 72 LLLAVVSL 79
 |||||
 Db 6 LLLAVVSL 13
 RESULT 70
 Q8QPL0 PRELIMINARY; PRT; 568 AA.
 ID Q8QPL0;
 AC Q8QPL0;
 DT 01-JUN-2002 (TrEMBLrel. 21, Created)
 DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Hemagglutinin (Fragment).

irus (A/Goose/Hong Kong/3014.8/2000 (H5N1)).
A negative-strand viruses; Orthomyxoviridae;
iruses; Influenzavirus A.
6675;

[N.A.
e/Hong Kong/3014.8/2000 (H5N1);

832; PubMed=11878904;
is M., Kong K.F., Dyrting K.C., Ellis T.M., Sit T.,
horridge K.F.;
za Viruses Isolated from Geese in Southeastern China:
Genetic Reassortment and Interspecies Transmission to

16-23 (2002).
HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
PTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
HOMOPRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
Y: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
12; AAL31388.1; -.
1; C: Viral envelope; IEA.
008980; Capsid hemag.
001364; Hemagglutn.
; Hemagglutinin; 1.
29; HEMAGGLUTININ2.
1225; Hemagglutn; 1.
ein; Glycoprotein; Hemagglutinin.
68 568
8 AA; 64281 MW; 0B0A4CFE034F1769 CRC64;

Query Match 2.8%; Score 8; DB 12; Length 568;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 0; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
VWSL 79
|||||
VWSL 13
PRELIMINARY; PRT; 620 AA.
TREMBLrel. 22, Created)
TREMBLrel. 22, Last sequence update)
TREMBLrel. 24, Last annotation update)
transporter.
06.
ampestris (pv. campestris).
teobacteria; Gammaproteobacteria; Xanthomonadales;
eae; Xanthomonas.
0;
[N.A.
3913 / NCPBB 528;
145; PubMed=1204217;
R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,
Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
Cannavan F., Cardozo J., Chambergo F., Ciapina L.P.,
I.B., Coutinho L.L., Cursino-Santos J.R., El-Dorri H.,
Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
I.F., Franco M.C., Greggio C.C., Gruber A.,
Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
T., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
Santos M., Truffi D., Tsai S.M., White F.F.,
Kitajima J.P.,
of the genomes of two Xanthomonas pathogens with differing

RT host specificities.";
RL Nature 417:459-463 (2002).
DR EMBL; AE012240; AAM40703.1; -.
GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0005215; P:transporter activity; IEA.
DR GO; GO:0006857; P:oligopeptide transport; IEA.
DR InterPro; IPR000109; PTR2.
DR Pfam; PF00854; PTR2; 1.
DR PROSITE; PS01022; PTR2_1; 1.
KW Complete proteome.
SQ SEQUENCE 620 AA; 67314 MW; E9904BFF039B6AEC CRC64;

Query Match 2.8%; Score 8; DB 16; Length 620;
Best Local Similarity 100.0%; Pred. No. 78;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 65 LALACLGL 72

Db 349 LALACLGL 356

RESULT 72

Q9N8H2

ID Q9N8H2 PRELIMINARY; PRT; 656 AA.

AC Q9N8H2

DT 01-OCT-2000 (TREMBlrel. 15, Created)

DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)

DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)

DE Hypothetical protein.

GN TB927.1.3840.

OS Trypanosoma brucei.

OC Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypano-

OX NCBI_TaxID=5691;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=TREU927;

RA Hall N., Berriman M., Lennard N.J., Harris B.R., Gerrard C.S.,

RA Atkin R.J., Barron A.J., Bart-Delabesse E.N., Bowman S.,

RA Bray-Allen S.P., Bringaund F., Clark L.N., Corton C.H., Cronin A.,

RA Davies R., Doggett J., Fraser A., Gruter E., Hall S., Harper D.A.

RA Hertz-Fowler C., Kay M.P., Leech V., Mayes R., Price C., Quail M.

RA Rabinowitsch E., Rutherford K., Sasse J., Sharp S., Showkhen R.

RA Gull K., Barrell B.G., Melville S.E.;

RT "The sequence and analysis of the highly polymorphic chromosome :

RT the African trypanosome, Trypanosoma brucei.";

RL Submitted (SRP-2002) to the EMBL/GenBank/DBJ databases.

DR EMBL; AL929607; CAB95571.1; -.

KW Hypothetical protein.

SQ SEQUENCE 656 AA; 72138 MW; CBAC892D25937FAD CRC64;

QY 93 BELVAEED 100

Db 454 BELVAEED 461

RESULT 73

Q9N8U8

ID Q9N8U8 PRELIMINARY; PRT; 1523 AA.

AC Q9N8U8

DT 01-OCT-2000 (TREMBlrel. 15, Created)

DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)

DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)

DE Hypothetical protein.

GN TB927.1.1600.

OS Trypanosoma brucei.

OC Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypano-

OX NCBI_TaxID=5691;

RN [1]

RP SEQUENCE FROM N.A.

GenCore version 5.1.6

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n search, using sw model

il 7, 2004, 17:47:28 ; Search time 60 Seconds
 (without alignments)
 1337.392 Million cell updates/sec

09-245-198A-4

SLDPEISARKLPLRSLG.....PWAHLKAAPFLTYGLFQVH 284

GO

op 60.0 , Gapext 60.0

6107 seqs, 282547505 residues

s satisfying chosen parameters: 1586107

th: 0

th: 2000000000

sting first 100 summaries

Genesep_29Jan04:*

genesep1980s:*

genesep1990s:*

genesep2000s:*

genesep2001s:*

genesep2002s:*

genesep2003as:*

genesep2003bs:*

genesep2004s:*

the number of results predicted by chance to have a
 than or equal to the score of the result being printed,
 d by analysis of the total score distribution.

SUMMARIES

Ch	Length	ID	Description
0	284	2	Aaw47525 Homo sapi
7	249	2	Aay09369 Human tum
7	249	3	Aay95338 Human PRO
7	249	3	Aab07526 Amino aci
7	249	5	Aau86129 Human PRO
7	249	6	Abr42315 Human TWE
7	249	7	Adc35206 Human TNF
9	249	7	Aaw29745 TNF relat
9	249	4	Aae00891 Human TRE
9	273	4	Aau03499 TWEAK ext
4	146	4	Aae00895 Human TRE
4	189	4	Aaw29746 TNF relat
4	189	4	Aae00892 Human UL4
7	208	2	Aaw93590 Human TNR
7	211	2	Aaw93591 Mouse TNR
3	225	2	Aaw47524 Mus muscu
3	225	3	Aab07527 Amino aci
3	249	7	Adc97712 Murine FL
12	58	3	Aag01265 Human sec
12	58	3	Aag01266 Human sec
12	365	6	Abra41235 Human DIT
12	748	2	Aay14906 Extended
12	749	5	Abb73512 M vaccae
18	54	4	Aau51863 Propionib
18	54	6	Abm48382 Propionib

Aam20858	Pej	55	2.8	8	26
Abb42637	Pej	55	2.8	8	27
Aam36451	Pej	55	2.8	8	28
Abb25983	Pr	55	2.8	8	29
Aam76342	Hu	55	2.8	8	30
Aam63528	Hu	55	2.8	8	31
Abg58050	Hu	55	2.8	8	32
Abg45635	Hu	55	2.8	8	33
Aam21621	Pej	65	2.8	8	34
Abb43981	Pej	65	2.8	8	35
Aam37923	Pej	65	2.8	8	36
Abb26890	Pr	65	2.8	8	37
Aam77706	Hu	65	2.8	8	38
Aam64984	Hu	65	2.8	8	39
Abg59361	Hu	65	2.8	8	40
Abg46737	Hu	65	2.8	8	41
Abg03723	No	69	2.8	8	42
Abg03663	No	71	2.8	8	43
Aau61282	Pr	84	2.8	8	44
Abm57801	Pr	84	2.8	8	45
Abg20259	No	110	2.8	8	46
Aao08094	Hu	117	2.8	8	47
Abb67712	Dr	184	2.8	8	48
Abp28041	SC	130	2.8	8	49
Aam23684	Hu	198	2.8	8	50
Abp41674	Hu	222	2.8	8	51
Abg16279	No	286	2.8	8	52
Abg01186	No	307	2.8	8	53
Abp79952	Rai	342	2.8	8	54
Abg13391	No	370	2.8	8	55
Abg05012	No	370	2.8	8	56
Abg18115	No	370	2.8	8	57
Abu36537	Pr	372	2.8	8	58
Abg15613	No	424	2.8	8	59
Aam23752	Hu	430	2.8	8	60
Adc08203	Ri	431	2.8	8	61
Adc64562	Sy	431	2.8	8	62
Abb57908	Dr	454	2.8	8	63
Ada54710	Hu	472	2.8	8	64
Abg20260	No	586	2.8	8	65
Ad46285	Hu	586	2.8	8	66
Ade62980	Hu	586	2.8	8	67
Aau32148	No	592	2.8	8	68
Abg03722	No	603	2.8	8	69
Ade79006	Hu	617	2.8	8	70
Abg20261	No	633	2.8	8	71
Abg28291	No	799	2.8	8	72
Ade08475	No	842	2.8	8	73
Abu16705	Pr	1032	2.8	8	74
Ada33851	Ac	1033	2.8	8	75
Aar66780	Ce	12	2.5	7	76
Abp82261	G I	16	2.5	7	77
Abb09088	Hu	17	2.5	7	78
Abg62026	Hu	18	2.5	7	79
Ada50609	HC	19	2.5	7	80
Adc99453	Car	20	2.5	7	81
Abg62030	Hu	23	2.5	7	82
Aay34189	Hu	24	2.5	7	83
Aab18622	An	25	2.5	7	84
Aaw89072	Pe	44	2.5	7	85
Aae01449	Hu	44	2.5	7	86
Aab51243	Hu	44	2.5	7	87
Abg63884	Hu	44	2.5	7	88
Abg63885	Hu	44	2.5	7	89
Abo45500	No	44	2.5	7	90
Abo26980	Pr	44	2.5	7	91
Abg02779	No	46	2.5	7	92
Aab40432	Hu	55	2.5	7	93
Aau63542	Pr	55	2.5	7	94
Abm60061	Pr	55	2.5	7	95
Abp00382	Hu	58	2.5	7	96
Aay12852	Hu	65	2.5	7	97
					98
					99

06:25:19 2004

us-09-245-198a-4.oligo.rag

2.5 66 7 ADC97100
2.5 69 4 ABB65500

ALIGNMENTS

idard; protein; 284 AA.

(first entry)

tumour necrosis factor related ligand (TRELL).

r necrosis factor related ligand; tnfr; treatment; cancer;
isease; immune system; stimulation; suppression;
ion.

97WO-US013945.

96US-0023541P.

96US-0028515P.

97US-0040820P.

EN INC.

GENEVA FACULTY MEDICINE.

he Y, Browning JL;

5619/13.

9600.

sis factor related ligand - useful for, e.g. treating cancer,
isease and immune responses to tissue grafts.

je 50-51; 69pp; English.

is that of human tumour necrosis factor related ligand
LL or active fragments can be included with a carrier in
al compositions to treat cancer, autoimmune diseases or
ases to tissue grafts, or to stimulate or suppress the immune
s useful to screen for TRELL receptors, by labelling with a
abel and screening compositions for binding. Agents
with TRELL-receptor binding can also be screened for, can
nistered, optionally with interferon- gamma, to induce cell
at, suppress or alter immune responses (especially involving
arcinoma cells) involving a signal pathway between TRELL and
. It's coding sequence can be used in gene therapy for TRELL-
rders in mammals (especially humans), e.g. tumours,
nd inflammatory diseases or inherited genetic disorders, by
into cells, and expressing, therapeutically effective amounts
e.g. a virus comprising a gene encoding TRELL. It may also
the preparation of prepare probes for screening
hetic DNAs for TRELL-encoding sequences and for antisense

AA;

100.0%; Score 284; DB 2; Length 284;

larity 100.0%; Pred. No. 2e-252;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LDFEISARRLPLRSLGSGGAVRQAQPPAPMAARRSQRRRGPGTALLVPLA 60

Db 1 MSLDLFEISARRLPLRSLGSGGAVRQAQPPAPMAARRSQRRRGPGTAL
QY 61 LGGLGALACLGLLAVVSLGSRASLSAQEPAQOEELVAEEDQDPSELNPOTESQD
Db 61 LGGLGALACLGLLAVVSLGSRASLSAQEPAQOEELVAEEDQDPSELNPOTESQD
QY 121 NLRVPRRSAPKGRKTRARRAJAAHYEVHPRPGDGAQAGVDTVSGWEARINS
Db 121 NLRVPRRSAPKGRKTRARRAJAAHYEVHPRPGDGAQAGVDTVSGWEARINS
QY 181 YNRQIGEFIVTRAGLYLYYCQVHFDEGKAVYIKLIDLLVDGLALRCLEFSATAA
Db 181 YNRQIGEFIVTRAGLYLYYCQVHFDEGKAVYIKLIDLLVDGLALRCLEFSATAA
QY 241 QLRLQVSGLLALRPGSSLRIRTLPAWHLKAAPFLTYFGLFQVH 284
Db 241 QLRLQVSGLLALRPGSSLRIRTLPAWHLKAAPFLTYFGLFQVH 284

RESULT 2

AAY09369

ID AAY09369 standard; protein; 249 AA.

XX AC AAY09369;

XX DT 15-JUL-1999 (first entry)

XX DE Human tumour necrosis factor Apo-3 ligand protein sequence.

XX KW Human; tumour necrosis factor; Apo-3 ligand; lymphotoxin; apopto

XX KW NF-kappaB-dependent transcription; JNK/SAPK-dependent response;

XX OS Homo sapiens.

XX PN WO919490-A1.

XX PD 22-APR-1999.

XX PF 09-OCT-1998; 98WO-US021407.

XX PR 10-OCT-1997; 97US-0062037P.

XX PR 17-DEC-1997; 97US-0069862P.

XX PA (GETH) GENENTECH INC.

XX PI Ashkenazi AJ, Marsters SA, Pitti R;

XX DR WPI; 1999-287982/24.

XX DR N-PSDB; AAX56000.

XX PT New human Apo3- ligand (a tumor necrosis factor) homologue.

XX PS Claim 1; Fig 1; 74pp; English.

XX CC The present sequence represents a human tumour necrosis factor (

CC lymphotoxin homologue designated Apo-3 ligand. Apo-3 ligand has

CC cytostatic activity. Apo-3 ligand can be used to induce apoptosi

CC mammalian cancer cells, to induce NF-kappaB-dependent transcript

CC to induce JNK/SAPK-dependent responses in mammalian cells

SQ Sequence 249 AA;

Query Match 87.7%; Score 249; DB 2; Length 249;

Best Local Similarity 100.0%; Pred. No. 2.6e-220;

Matches 249; Conservative 0; Mismatches 0; Indels 0;

QY 36 MAARRSQRRRGPGTALLVPLALGGLGALACLGLLAVVSLGSRASLSAQEP.

Db 1 MAARRSQRRRGPGTALLVPLALGGLGALACLGLLAVVSLGSRASLSAQEP.

QY 96 VAEEDQDPSELNPOTESQDPAPFLNRLVPRRSAPKGRKTRARRAJAAHYEVH.

Db 61 VAEEDQDPSELNPOTESQDPAPFLNRLVPRRSAPKGRKTRARRAJAAHYEVH.

WDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYLYCQVHFDEGKAVYLKLD 215
 |||||
 WDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYLYCQVHFDEGKAVYLKLD 180
 |||||
 WDLALCLREFSNTAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWLKKAAPFL 275
 |||||
 WDLALCLREFSNTAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWLKKAAPFL 240

.FQVH 284
 |||||
 .FQVH 249

hard; protein; 249 AA.

(first entry)

antitumour protein.

antitumour; tumour; therapy; cytostatic; breast cancer;
 ; renal cancer; colorectal cancer; uterine cancer;
 ; lung cancer; bladder cancer;
 is system cancer; melanoma; leukaemia; neoplasm.

Location/Qualifiers

1. .40
 /label= Signal_peptide
 10. .14
 /note= "amidation"
 24. .35
 /note= "prokaryotic membrane lipoprotein lipid"
 27. .33
 /note= "N-myristoylation"
 29. .35
 /note= "N-myristoylation"
 36. .42
 /note= "N-myristoylation"
 41. .249
 /label= PRO207
 45. .51
 /note= "N-myristoylation"
 97. .101
 /note= "amidation"
 118. .124
 /note= "N-myristoylation"
 121. .127
 /note= "N-myristoylation"
 125. .131
 /note= "N-myristoylation"
 128. .134
 /note= "N-myristoylation"
 139. .143
 /note= "Asn is N-glycosylated"

2.

99WO-US028565.
 98US-0113296P.
 99WO-US005028.
 99US-0130232P.
 99US-0131445P.
 99US-0134287P.
 99US-0144758P.

PR 26-JUL-1999; 99US-0145698P.
 PR 15-SEP-1999; 99WO-US021090.
 PR 15-SEP-1999; 99WO-US021547.
 XX
 PA (GETH) GENENTECH INC.
 XX Ashkenazi AJ, Goddard A, Godowski PJ, Gurney AL, Marsters SA;
 PI Napier MA, Pitti RM, Wood WI;
 XX WPI; 2000-442668/38.
 DR N-PSDB; AAA49717.
 XX
 PT Novel composition to inhibit neoplastic cell growth or for treati
 in mammal comprises polypeptides PRO179, PRO207, PRO320, PRO219,
 PT PRO224, PRO328, PRO301, PRO526, PRO362, PRO356, PRO509 or PRO866.
 XX
 PS Claim 19; Fig 4; 172pp; English.
 XX
 CC The present sequence is that of human antitumour protein PRO207,
 deduced from a foetal kidney cDNA clone (see AAA49717). PRO207 sh
 CC amino acid sequence identity to tumour necrosis factor mem
 CC especially human lymphotoxin-beta (23.4%) and human CD40 ligand (
 CC Mol.wt. is 27.216. A claimed method for inhibiting the growth of
 CC cell comprises exposing the tumor cell to PRO179, PRO207, PRO320,
 CC PRO221, PRO224, PRO328, PRO301, PRO526, PRO362, PRO356, PRO509 or
 CC (see AA95337-49), their agonists or chimeric polypeptides incorp
 CC them. The tumour is especially a cancer selected from breast, ova
 CC renal, colorectal, uterine, prostate, lung, bladder and central n
 CC system cancer, melanoma and leukaemia. Methods for the recombinan
 CC expression of the antitumour proteins are also provided

XX Sequence 249 AA;

Query Match 87.7%; Score 249; DB 3; Length 249;
 Best Local Similarity 100.0%; Pred. No. 2.6e-220;
 Matches 249; Conservative 0; Mismatches 0; Indels 0; G

QY 36 MAARRSQRRGRGEGPTALLVPLALGLALCLGLLLAVVSLGSRASLSAQEPA
 |||||
 Db 1 MAARRSQRRGRGEGPTALLVPLALGLALCLGLLLAVVSLGSRASLSAQEPA
 |||||
 QY 96 VAEEDQDPSELNPQTEESQDPAPFLNLRVPRSPKGRKTRARRAIAAHYEVHPR
 |||||
 Db 61 VAEEDQDPSELNPQTEESQDPAPFLNLRVPRSPKGRKTRARRAIAAHYEVHPR
 |||||
 QY 156 GAQAGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYLYCQVHFDEGKAVY
 |||||
 Db 121 GAQAGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYLYCQVHFDEGKAVY
 |||||
 QY 216 LLVDGVLALRCLREFSNTAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWLKKA
 |||||
 Db 181 LLVDGVLALRCLREFSNTAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWLKKA
 |||||
 QY 276 TYFGLFQVH 284
 |||||
 Db 241 TYFGLFQVH 249

RESULT 4

AAB07526
 ID AAB07526 standard; protein; 249 AA.
 XX
 AC AAB07526;
 XX
 XX 20-OCT-2000 (first entry)

XX Amino acid sequence of a soluble recombinant human TWEAK protein.
 DE TWEAK protein; immunological disorder; immune response; inflammat
 XX TWEAK blocking agent; autoimmune disease; organ transplant reject
 KW Graft-versus-Host disease; GVHD; lymphoid cell malignancy; shock;
 XX Homo sapiens.
 OS

AI.

2000WO-US001044.

99US-0116168P.

EN INC.

6036/41.

nd treating immune responses using modulators, especially of TWEAK, TWEAK receptors and TWEAK ligands, useful for inflammation and graft versus host disease.

Fig 1; 45pp; English.

sequence represents a TWEAK protein. The specification method for preventing or treating an immunological disorder using an immune response in an animal. The method comprises a TWEAK blocking agent. The method may be used for nd treating immune disorders associated with inappropriate nd/or activity of TWEAK. These disorders include autoimmune and chronic inflammation, organ transplant rejection, -Host disease (GVHD), lymphoid cell malignancies, septic and of shock, loss of immune responsiveness (as seen in human ency virus (HIV) infections) and failure of the immune tumour growth

AA;

87.7%; Score 249; DB 3; Length 249; larity 100.0%; Pred. No. 2.6e-220; Indels 0; Gaps 0; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RRSQRRRGRRGPGTALLVPLALGLGALACGLLLAVSLGSRASLSAQEEL 95

RRSQRRRGRRGPGTALLVPLALGLGALACGLLLAVSLGSRASLSAQEEL 60

EDQDPSLNQTESQDPAPFLNLRVPRRSAPKGRKTRARRAIAAHYVHPRGD 155

EDQDPSLNQTESQDPAPFLNLRVPRRSAPKGRKTRARRAIAAHYVHPRGD 120

AGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYCQVHDEGKAVYKLD 215

AGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYCQVHDEGKAVYKLD 180

DGVLAIRCLFEFSATAASSIGPOLRLCQVSGLLALRPGSSLRITLTPWAHLKAAPFL 275

DGVLAIRCLFEFSATAASSIGPOLRLCQVSGLLALRPGSSLRITLTPWAHLKAAPFL 240

3LFQVH 284

3LFQVH 249

adard; protein; 249 AA.

(first entry)

polypeptide.

benign tumour; malignant tumour; lymphoid malignancy; autonal disorder; stromal disorder; blastocoeic disorder; disorder; immune disorder; angiogenic disorder; cytostatic;

KW neuroprotective.

XX Homo sapiens.

OS WO200153486-A1.

XX 26-JUL-2001.

XX 11-FEB-2000; 2000WO-US003565.

XX 08-MAR-1999; 99WO-US005028.

XX 11-MAR-1999; 99US-0123972P.

XX 11-MAY-1999; 99US-0133459P.

XX 02-JUN-1999; 99WO-US012252.

XX 22-JUN-1999; 99US-0140650P.

XX 22-JUN-1999; 99US-0140653P.

XX 20-JUL-1999; 99US-0144758P.

XX 26-JUL-1999; 99US-0145698P.

XX 28-JUL-1999; 99US-0146222P.

XX 17-AUG-1999; 99US-0149395P.

XX 31-AUG-1999; 99US-0151689P.

XX 01-SEP-1999; 99WO-US020111.

XX 15-SEP-1999; 99WO-US021090.

XX 30-NOV-1999; 99WO-US028313.

XX 01-DEC-1999; 99WO-US028301.

XX 01-DEC-1999; 99WO-US028634.

XX 05-JAN-2000; 2000WO-US000219.

(GETH) GENENTECH INC.

XX Ashkenazi AJ, Goddard A, Godowski PJ, Gurney AL, Hillan KJ;

XX Marsters SA, Pan J, Pitti RM, Roy MA, Smith V, Stone DM;

XX Watanabe CK, Wood WI;

XX WPI: 2002-205567/26.

XX N-PSDB; ABK40255.

XX Thirty five nucleic acids encoding PRO polypeptides, useful for benign or malignant tumors, leukemias and lymphoid malignancies, inflammatory, angiogenic and immunologic disorders.

XX Claim 61; Fig 4; 302pp; English.

XX The present invention relates to the isolation of novel human PR polypeptides and the polynucleotide sequences encoding them. The polypeptides, agonists, antagonists or anti-PRO antibodies are u treating benign or malignant tumors (e.g. renal, kidney, bladder breast, etc), leukaemias and lymphoid malignancies, other disord as neuronal, glial, astrocytal, hypothalamic, glandular, macroph stromal and blastocoeic disorders, inflammatory, immune and ang disorders. The polynucleotide sequences are also useful in gene disorders. The polynucleotide sequences are also useful in gene AAU86128-AAU86162 represent the human PRO polypeptides of the in

XX Sequence 249 AA;

Query Match 87.7%; Score 249; DB 5; Length 249;

Best Local Similarity 100.0%; Pred. No. 2.6e-220;

Matches 249; Conservative 0; Mismatches 0; Indels 0;

QY 36 MAARRSQRRRGRRGPGTALLVPLALGLGALACGLLLAVSLGSRASLSAQEP.

Db 1 MAARRSQRRRGRRGPGTALLVPLALGLGALACGLLLAVSLGSRASLSAQEP.

QY 96 VAEEDQDPSLNQTESQDPAPFLNLRVPRRSAPKGRKTRARRAIAAHYVHP.

Db 61 VAEEDQDPSLNQTESQDPAPFLNLRVPRRSAPKGRKTRARRAIAAHYVHP.

QY 156 GAQAGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYCQVHDEGKAV

Db 121 GAQAGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYCQVHDEGKAV

QY 216 LIAVDGVLALRCLFEFSATAASSIGPOLRLCQVSGLLALRPGSSLRITLTPWAHLK

QVLAALRCLEEFSAATAASSLGPQLRLCQVSLALRPQSSLRIRTLPLWAHLK 240

LFQVH 284

LFQVH 249

ard; protein; 249 AA.

(first entry)

rotein.

tumour necrosis factor; ligand; cytostatic;
or; osteopathic.

A2.

2002WO-US023782.

2001US-0307838P.

GENOME SCI INC.

Rosen CA;

659/40.

901.

timeric complex having a first polypeptide member of the
s factor (TNF) ligand family, and a second different member
family, useful for treating cancer, osteoporosis or an
sease.

age 368-369; 388pp; English.

equence is the protein sequence for human TWEAK protein. The
ates to compositions comprising heterotrimeric complexes of
is factor (TNF) ligand family members, and their use in the
event and treatment of disease. In one embodiment, the
c complex comprises full-length or extracellular portions of
1-length or extracellular portions of other TNF ligand
s, preferably VEGI or VEGI-SV. The heterotrimeric complexes
ion are useful for treating an autoimmune disease, cancer or
and particularly for inhibiting cancer cell proliferation,
cell proliferation, or inducing apoptosis of T cells

AA;

87.7%; Score 249; DB 6; Length 249;

arity 100.0%; Pred. No. 2.6e-220;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

RSQRRRGRRGPGTALLVPLALGLALACLGILLAVVSLGSRASLSAQEPQEEEL 95

RSQRRRGRRGPGTALLVPLALGLALACLGILLAVVSLGSRASLSAQEPQEEEL 60

QDPSELNPQTESQDPAPFLNLRPRRSPKGRKTRARRAJAAHYEVHPRPGQD 155

QDPSELNPQTESQDPAPFLNLRPRRSPKGRKTRARRAJAAHYEVHPRPGQD 120

3VDGTSGWEARINSSSPRYNRQIGEFIVTRAGLYLYLCQVHDEGKAVYKLD 215

3VDGTSGWEARINSSSPRYNRQIGEFIVTRAGLYLYLCQVHDEGKAVYKLD 180

QY 216 LLVDGVLAALRCLEEFSAATAASSLGPQLRLCQVSLALRPQSSLRIRTLPLWAHLK

Db 181 LLVDGVLAALRCLEEFSAATAASSLGPQLRLCQVSLALRPQSSLRIRTLPLWAHLK

QY 276 TYFGLFQVH 284

Db 241 TYFGLFQVH 249

RESULT 7

ADC35206

ID ADC35206 standard; protein; 249 AA.

XX AC ADC35206;

DT 18-DEC-2003 (first entry)

XX Human TNF ligand family member #12.

XX human; tumour necrosis factor; TNF ligand; endokine alpha;
XX excessive bone resorption disorder; osteoporosis; Paget's disease;
XX arterial calcification.

OS Homo sapiens.

PN US2003100074-A1.

PD 29-MAY-2003.

XX 15-AUG-2002; 2002US-00218547.

PR 16-AUG-2001; 2001US-0312542P.

PR 30-OCT-2001; 2001US-0330761P.

XX (YUGG/) YU G.

PA (NIJJ/) NI J.

PA (ROSE/) ROSEN C A.

PA (NARD/) NARDELLI B.

XX Yu G, Ni J, Rosen CA, Nardelli B;

DR WPI; 2003-696072/66.

DR N-PSDB; ADC35205.

XX New Endokine alpha gene useful for preparing a composition for tr
disease associated with excessive or insufficient bone resorptior
osteoporosis, Paget's disease or arterial calcification.

XX Disclosure; SEQ ID NO 24; 145pp; English.

CC The invention relates to an isolated nucleic acid molecule encodi
tumour necrosis factor family ligand. A composition comprising th
isolated antibody or its fragment is used for treating an individ
need of decreased level of endokine alpha activity. The endokine
polypeptide present in a heterotrimeric complex is used for treat
individual having a disorder associated with excessive bone resor
e.g. osteoporosis, Paget's disease or arterial calcification. The
individual having a disorder associated with insufficient bone re
comprises administering an endokine alpha antagonist, which is th
antibody that binds specifically to endokine alpha polypeptide. T
CC present sequence represents the amino acid sequence of a tumour n
factor family ligand.

XX Sequence 249 AA;

Query Match

Best Local Similarity 87.7%; Score 249; DB 7; Length 249;

Matches 249; Conservative 0; Mismatches 0; Indels 0; G

QY 36 MAARRSQRRRRGRRGPGTALLVPLALGLALACLGILLAVVSLGSRASLSAQEPA

Db 1 MAARRSQRRRRGRRGPGTALLVPLALGLALACLGILLAVVSLGSRASLSAQEPA

06:25:19 2004

us-09-245-198a-4.oligo.rag

EDQPSLNQTESQDPAPFLNRLVPRRSAPKGRKTRARRAJAAHYEVHPRGQD 155
|||||
EDQPSLNQTESQDPAPFLNRLVPRRSAPKGRKTRARRAJAAHYEVHPRGQD 120
|||||
AGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYYCQVHFDGKAVYKLD 215
|||||
AGVDGTVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYYCQVHFDGKAVYKLD 180
|||||
DGVLAIRCLBEFSATAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWHLKAAPFL 275
|||||
DGVLAIRCLBEFSATAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWHLKAAPFL 240
|||||
3LFOVH 284
|||||
3LFOVH 249
|||||
ndard; protein; 249 AA.

(first entry)
endothelium proliferative agent protein.
lium proliferative agent; TREPA; wound healing; cancer;
ing; vascularisation; apoptosis; autoimmune; birth control.

98WO-US002859.
97US-00798692.
98US-00021706.
IT LAB.

7255/38.
7613.
cleic acid encoding TREPA - useful for diagnosis and
autoimmune disease, tumours and inflammation.
3e 123-4; 142pp; English.
ted endothelium proliferative agent (TREPA), or its
agonists, are used to treat a deficit of TREPA, e.g. to
healing or tissue grafting, by promoting vascularisation,
ce apoptosis for treating cancer and eliminating autoreactive
an adjunct to cancer chemotherapy or antiviral treatment.
es can also be used to target cytotoxic agents or for
lation of the corresponding receptor, the nucleic acid for
used to transform tumour cells to render them more
o TREPA and to screen for TREPA mimics. Ribozymes, antisense
dies or peptides, are used to treat TREPA-associated
g. tumours and metastases (by inhibiting vascularisation),
or a wide range of autoimmune conditions, conditions
normal stimulation of epithelial cells (e.g.
sis), for birth control (inhibiting ovulation and placental
r other angiogenic conditions (e.g. ulcers)
AA;
84.9%; Score 241; DB 2; Length 249;
larity 100.0%; Pred. No. 5.9e-213;

Matches 241; Conservative 0; Mismatches 0; Indels 0;
QY 44 RRRRGCEPTALLVPLALGGLALACLGALLAVVSLGSRASLSAQEPAGEELVAE
|||||
Db 9 RRRRGCEPTALLVPLALGGLALACLGALLAVVSLGSRASLSAQEPAGEELVAE
|||||
QY 104 SELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAJAAHYEVHPRGQDGAQ
|||||
Db 69 SELNPQTESQDPAPFLNRLVPRRSAPKGRKTRARRAJAAHYEVHPRGQDGAQ
|||||
QY 164 TVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYYCQVHFDGKAVYKLDLLV
|||||
Db 129 TVSGWEEARINSSPLRYNRQIGEFIVTRAGLYLYYCQVHFDGKAVYKLDLLV
|||||
QY 224 LRCLEFSATAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWHLKAAPFLTYF
|||||
Db 189 LRCLEFSATAASSLGPQLRLCQVSGLLALRPGSSLRIRTLPAWHLKAAPFLTYF
|||||
QY 284 H 284
|||||
Db 249 H 249
|||||
RESULT 9
AAE00891
ID AAE00891 standard; protein; 249 AA.
XX
AC AAE00891;
XX
DT 04-JUL-2001 (first entry)
XX
DE Human TREPA (TNF related endothelium proliferative agent).
XX
KW Human; tumour necrosis factor; TNF; angiogenesis; wound healing;
KW TNF related endothelium proliferative agent; tumour; metastasis;
KW grafting; vulneryary.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Domain 98..249
FT /label= Extracellular_domain
XX
PN US6207642-B1.
XX
PD 27-MAR-2001.
XX
PF 26-JUN-1998; 98US-00105343.
XX
PR 12-FEB-1997; 97US-00798692.
PR 10-FEB-1998; 98US-00021706.
XX
PA (ABBO) ABBOTT LAB.
XX
PI Wiley SR;
XX
DR WPI; 2001-280760/29.
DR N-PSDB; AAD04350.
XX
PT Inducing angiogenesis in mammal at desired sites for promoting w
PT healing, by administering soluble fragment of extracellular doma
PT tumor necrosis factor related endothelium proliferative agent pr
XX
PS Claim 1; Col 75-76; 53pp; English.
XX
CC The present invention relates to extracellular signal molecules,
CC particularly members of tumour necrosis factor (TNF) family mole
CC designated as TREPA (TNF related endothelium proliferative agent
CC Soluble biologically active TREPA are used to treat TREPA-associ
CC diseases, tumours or metastases. TREPA is used for inducing angi
CC in human for promoting wound healing and for vascularising graft
CC for successful grafting and to promote tissue grafts. The presen
CC acid sequence is clone ID #690050 human TREPA

A: 84.9%; Score 241; DB 4; Length 249;
 rity 100.0%; Pred. No. 5.9e-213; Indels 0; Gaps 0;
 nservative 0; Mismatches 0; Indels 0; Gaps 0;
 GEPGTALLVPLALGLGLALACIGLLAVVSLGSRASLSAQEPQAEELVAEEDQDP 103
 GEPGTALLVPLALGLGLALACIGLLAVVSLGSRASLSAQEPQAEELVAEEDQDP 68
 PTEESQDPAPFLNRLVRRSAPKGRKTRARRAIAAHYEVHPRPQDGAQAGVDG 163
 PTEESQDPAPFLNRLVRRSAPKGRKTRARRAIAAHYEVHPRPQDGAQAGVDG 128
 TEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVYKLDLLVDGVLA 223
 TEARINSSPLRYNQIGEFIVTRAGLYLYCQVHFDEGKAVYKLDLLVDGVLA 198
 EFSATAASLSGQPLRLCOVSGLLALRPQSSLRIRTLPAWHLKAAPFLTYFGLFQV 283
 EFSATAASLSGQPLRLCOVSGLLALRPQSSLRIRTLPAWHLKAAPFLTYFGLFQV 248

ard; protein; 273 AA.

first entry)

lular domain-containing fusion protein.

lular domain; tumour necrosis factor; TNF; angiogenesis;
 ularisation; diabetic retinopathy; neovascular glaucoma;
 ; retinopathy of prematurity; retrolental fibroplasia;
 tis; macular degeneration; arthritis; rheumatism;
 neovascularisation; psoriasis; metastatic condition;
 ur; sarcoma; carcinoma; benign tumour; haemophilic joint;
 condition; myocardial angiogenesis; wound granulation;
 ascular adhesion; telangiectasia; ischaemia; human;
 c plaque neovascularisation; coronary atherosclerosis;
 erosclerosis; PDC409-LZ-TWEAK; TWEAK receptor; TWEAKR;

000WO-U5034755.

99US-0172878P.

000US-0203347P.

X CORP.

75/44.

64.

ogenesis in a mammal for treating diseases mediated by
 e.g. solid tumors and vascular deficiencies of cardiac or
 sue, by administering antagonist or agonist of TWEAK

XX The sequence represents a fusion protein encoded by the express
 PS vector pDC409-LZ-TWEAK. The fusion protein comprises a growth hor
 CC leader, a leucine zipper multimerisation domain, and the extracel
 CC domain of human TWEAK. The fusion protein was used in the isolati
 CC human TWEAK receptor (TWEAKR)-expressing clones from a COS cell h
 CC cDNA library. The TWEAK protein is a member of the tumour necrosi
 CC (TNF) family and induces angiogenesis. TWEAKR may therefore be us
 CC screen for and develop TWEAK agonists and antagonists for the mo
 CC of angiogenesis, to be used in the treatment and diagnosis of hum
 CC disease. The disorders mediated by angiogenesis include ocular di
 CC characterised by ocular neovascularisation such as diabetic retin
 CC neovascular glaucoma, retinoblastoma, retinopathy of prematurity,
 CC retrolental fibroplasia, rubeosis, uveitis, macular degeneration,
 CC corneal graft neovascularisation, and inflammatory diseases such
 CC arthritis, rheumatism and psoriasis. Other treatable diseases inc
 CC malignant and metastatic conditions such as sarcomas and carcinom
 CC benign tumours and preneoplastic conditions, myocardial angiogen
 CC haemophilic joints, scleroderma, vascular adhesions, atherosclero
 CC plaque neovascularisation, telangiectasia, wound granulation, cor
 CC atherosclerosis, peripheral atherosclerosis and ischaemia

XX Sequence 273 AA;

Query Match 72.9%; Score 207; DB 4; Length 273;
 Best Local Similarity 100.0%; Pred. No. 1.2e-181;
 Matches 207; Conservative 0; Mismatches 0; Indels 0; G;

QY 78 SLGSRASLSAQEPQAEELVAEEDQDPSELNPQTESQDPAPFLNRLVRRSAPKGI

Db 67 SLGSRASLSAQEPQAEELVAEEDQDPSELNPQTESQDPAPFLNRLVRRSAPKGI

QY 138 ARRAIAAHYEVHPRPQDGAQAGVDGTVSGWEARINSSPLRYNQIGEFIVTRA

Db 127 ARRAIAAHYEVHPRPQDGAQAGVDGTVSGWEARINSSPLRYNQIGEFIVTRA

QY 198 LYCQVHFDEGKAVYKLDLLVDGVLAALRCLEFFSATAASSLGQPLRLCOVSGLLAL

Db 187 LYCQVHFDEGKAVYKLDLLVDGVLAALRCLEFFSATAASSLGQPLRLCOVSGLLAL

QY 258 SLRIRTLPAWHLKAAPFLTYFGLFQVH 284

Db 247 SLRIRTLPAWHLKAAPFLTYFGLFQVH 273

RESULT 11

AAE00895

ID AAE00895 standard; protein; 146 AA.

XX AAE00895;

XX 04-JUL-2001 (first entry)

DE Human TREPA (TNF related endothelium proliferative agent) fragment

KW Human; tumour necrosis factor; TNF; angiogenesis; wound healing; 1

KW TNF related endothelium proliferative agent; tumour; metastasis;

XX grafting; vulnery.

OS Homo sapiens.

PN US6207642-B1.

XX 27-MAR-2001.

XX 26-JUN-1998; 98US-00105343.

XX 12-FEB-1997; 97US-00798692.

PR 10-FEB-1998; 98US-00021706.

XX (ABCO) ABBOTT LAB.

XX
 CC The TNF-related endothelium proliferative agent (TREPA), or its
 CC activators or agonists, are used to treat a deficit of TREPA, e.
 CC promote wound healing or tissue grafting, by promoting vasculari-
 CC also to induce apoptosis for treating cancer and eliminating aut
 CC T cells, as an adjunct to cancer chemotherapy or antiviral treat
 CC TREPA peptides can also be used to target cytotoxic agents or fo
 CC affinity isolation of the corresponding receptor, the nucleic ac
 CC which can be used to transform tumour cells to render them more
 CC responsive to TREPA and to screen for TREPA mimics. Ribozymes, a
 CC RNA, antibodies or peptides, are used to treat TREPA-associated
 CC diseases, e.g. tumours and metastases (by inhibiting vascularisa
 CC inflammation or a wide range of autoimmune conditions, condition
 CC involving abnormal stimulation of epithelial cells (e.g.
 CC atherosclerosis), for birth control (inhibiting ovulation and pl
 CC formation) or other angiogenic conditions (e.g. ulcers)
 XX
 SQ Sequence 189 AA;

Query Match 50.4%; Score 143; DB 2; Length 189;
 Best Local Similarity 100.0%; Pred. No. 5.7e-123;
 Matches 143; Conservative 0; Mismatches 0; Indels 0;
 QY 142 IAAHYEVHPRPGDGAQAGVDGTVSGWEARINSSPLRYNROI GEFIVTRAGLY
 Db 47 IAAHYEVHPRPGDGAQAGVDGTVSGWEARINSSPLRYNROI GEFIVTRAGLY
 QY 202 VHFDEGKAVYKLDLLVDGVLALRCLEEFSAATAASSLGPGQLRCQVSGLLALRPG
 Db 107 VHFDEGKAVYKLDLLVDGVLALRCLEEFSAATAASSLGPGQLRCQVSGLLALRPG
 QY 262 RTLPWAHLKAAPFLTYFGLFQVH 284
 Db 167 RTLPWAHLKAAPFLTYFGLFQVH 189

RESULT 13

AAE00892
 ID AAE00892 standard; protein; 189 AA.
 AC AAE00892;
 XX
 DT 04-JUL-2001 (first entry)
 XX
 DE Human UL4flag TREPA soluble construct.
 XX
 KW Human; tumour necrosis factor; TNF; angiogenesis; wound healing;
 KW TREPA; TNF related endothelium proliferative agent; metastasis;
 KW vulnery; HUVEC; human umbilical vein endothelial cell; UL4flag
 XX
 OS Homo sapiens.
 XX
 PN US6207642-B1.
 XX
 PD 27-MAR-2001.
 XX
 PF 26-JUN-1998; 98US-00105343.
 XX
 PR 12-FEB-1997; 97US-00798692.
 PR 10-FEB-1998; 98US-00021706.
 XX
 PA (ABBO) ABBOTT LAB.
 XX
 PI Wiley SR;
 XX
 DR WPI; 2001-280760/29.
 XX
 PT Inducing angiogenesis in mammal at desired sites for promoting w
 PT healing, by administering soluble fragment of extracellular doma
 PT tumor necrosis factor related endothelium proliferative agent pr
 XX
 PS Example 2; Col 75-78; 53pp; English.
 XX

AA;
 51.4%; Score 146; DB 4; Length 146;
 larity 100.0%; Pred. No. 7.8e-126; Mismatches 0; Indels 0; Gaps 0;
 Conservative 0; Mismatches 0; Indels 0;
 QY 142 IAAHYEVHPRPGDGAQAGVDGTVSGWEARINSSPLRYNROI GEFIVTRAGLYYL 198
 Db 47 IAAHYEVHPRPGDGAQAGVDGTVSGWEARINSSPLRYNROI GEFIVTRAGLYYL 60
 QY 202 VHFDEGKAVYKLDLLVDGVLALRCLEEFSAATAASSLGPGQLRCQVSGLLALRPGSS 258
 Db 107 VHFDEGKAVYKLDLLVDGVLALRCLEEFSAATAASSLGPGQLRCQVSGLLALRPGSS 120
 QY 262 RTLPWAHLKAAPFLTYFGLFQVH 284
 Db 167 RTLPWAHLKAAPFLTYFGLFQVH 146

idard; protein; 189 AA.

(first entry)

endothelium proliferative agent protein 2.

ium proliferative agent; TREPA; wound healing; cancer;
 ing; vascularisation; apoptosis; autoimmune; birth control.

98WO-US002859.

97US-00798692.

98US-00021706.

IT LAB.

7255/38.

leic acid encoding TREPA - useful for diagnosis and
 autoimmune disease, tumours and inflammation.

je 125-6; 142pp; English.

vention relates to extracellular signal molecules, members of tumour necrosis factor (TNF) family molecules (TREPA (TNF related endothelium proliferative agent)). Specially active TREPA are used to treat TREPA-associated tumors or metastases. TREPA is used for inducing angiogenesis promoting wound healing and for vascularising grafted tissue and grafting and to promote tissue grafts. The present amino acid sequence is human U4fag TREPA soluble construct. This sequence is biologically active molecule is capable of inducing HUVEC (human umbilical vein endothelial cells) cells

AA;
arity 50.4%; Score 143; DB 4; Length 189;
conservative 0; Mismatches 0; Indels 0; Gaps 0;

YEHPFGQGAQAGVDGTVSGWEAFARINSSPLRYNRQIGFIVTRAGLYLYYCQ 201
|||||
YEHPFGQGAQAGVDGTVSGWEAFARINSSPLRYNRQIGFIVTRAGLYLYYCQ 106
EGKAVYLKDLVDGVLALRCLEEFSAATASSLGQPLRLCQVSGLLALRPGSSLRI 261
EGKAVYLKDLVDGVLALRCLEEFSAATASSLGQPLRLCQVSGLLALRPGSSLRI 166

WAHLKAAPFLTYFGLFQVH 284
|||||
WAHLKAAPFLTYFGLFQVH 189

gird; protein; 208 AA.

(first entry)

rotein.

is factor receptor; signal transducer molecule; TNF; APO4; abnormality; gestational abnormality; prostate cancer; PO3; TNRL-1; TNRL-3; diagnosis; treatment; therapy; disease; main; immunogen; antibody preparation; breast carcinoma; nan.

98WO-US018393.

97US-00924634.

WASHINGTON.

191/17.
124.

rosis Factor family receptor polypeptides and ligands - agnosis and treatment of prostate cancer and developmental abnormalities.

13A; 156pp; English.

a describes isolated Tumor Necrosis Factor (TNF) family peptides: APO4, APO6, APO8 and APO9 or their active fragments. APO4 is useful for diagnosing prostate cancer by

CC determining levels of APO4 in an individual. Prostate cancer can
CC treated using APO4 selective binding agents linked to a therapeutic
CC moiety. APO4 polypeptides are also useful for identifying select-
CC binding agents, useful in diagnosis/treatment of disease by bind-
CC agents to the polypeptide/active fragment which is extracellular,
CC expressed on the cell surface. The binding is preferably performed
CC vivo. APO4 polypeptides/ active fragments are also useful for sci-
CC for agonists and antagonists by binding and observing the change;
CC activity. Effective pharmacological agents useful in diagnosis of
CC treatment of disease are also identified using APO4 polypeptides/
CC fragments and APO4 signal transducer molecules that specifically
CC with a cytoplasmic domain of APO4 and detecting a change in level
CC activity. The method is performed in vivo or in vitro. APO polype-
CC are all useful as immunogens for preparing antibodies. APO4 is al-
CC useful for diagnosis/treatment of developmental or gestational
CC abnormalities. APO8 was transfected to human breast carcinoma cell
CC MCF-7, and induced apoptosis

XX Sequence 208 AA;

Query Match 37.7%; Score 107; DB 2; Length 208;
Best Local Similarity 99.5%; Pred. No. 7.8e-90;
Matches 207; Conservative 0; Mismatches 1; Indels 0; C

Qy 77 VSLGSRASLSAQEPAQBELVAEEDQDPSELNPQTEESQDPAPFLNRLVRRPSAPP
Db 1 VSLGSRASLSAQEPAQBELVAEEDQDPSELNPQTEESQDPAPFLNRLVRRPSAPP
Qy 137 RARRAIAAHVEHPRPGQDGAQAGVDGTVSGWEAFARINSSPLRYNRQIGFIVTR
Db 61 RARRAIAAHVEHPRPGQDGAQAGVDGTVSGWEAFARINSSPLRYNRQIGFIVTR
Qy 197 YLYCOVHFDEGKAVYLKDLVDGVLALRCLEEFSAATASSLGQPLRLCQVSGLLP
Db 121 YLYCOVHFDEGKAVYLKDLVDGVLALRCLEEFSAATASSLGQPLRLCQVSGLLP
Qy 257 SSLRIRTLPPWAHLKAAPFLTYFGLFQVH 284
Db 181 SSLRIRTLPPWAHLKAAPFLTYFGLFQVH 208

RESULT 15
AAW93591
ID AAW93591 standard; protein; 211 AA.

AC AAW93591;

DT 18-JUN-1999 (first entry)

DE Mouse TNRL3 protein.

XX Tumour necrosis factor receptor; signal transducer molecule; TNF;
KW developmental abnormality; gestational abnormality; prostate ca-
KW APO6; APO8; APO9; TNRL-1; TNRL-3; diagnosis; treatment; therapy;
KW cytoplasmic domain; immunogen; antibody preparation; breast carci-
KW apoptosis; mouse.

OS Mus sp.

XX WO9911791-A2.

XX 11-MAR-1999.

XX 04-SEP-1998; 98WO-US018393.

XX 05-SEP-1997; 97US-00924634.

XX (UNIW) UNIV WASHINGTON.

XX Chaudhary PM;

XX WPI; 1999-205191/17.

DR N-PSDB; AAX23425.

rosis Factor family receptor polypeptides and ligands -
agnosis and treatment of prostate cancer and developmental
al abnormalities.

g 13B; 156pp; English.

on describes isolated Tumor Necrosis Factor (TNF) family
ypeptides: APO4, APO6, APO8 and APO9 or their active
nd isolated TNF related ligands 1 and 3 (TNRL1 and TNRL3) or
ragments. APO4 is useful for diagnosing prostate cancer by
levels of APO4 in an individual. Prostate cancer can also be
g APO4 selective binding agents linked to a therapeutic
polypeptides are also useful for identifying selective
ts, useful in diagnosis/treatment of disease by binding of
e polypeptide/active fragment which is extracellular, or
the cell surface. The binding is preferably performed in
olypeptides/ active fragments are also useful for screening
and antagonists by binding and observing the change in APO4
fective pharmacological agents useful in diagnosis or
disease are also identified using APO4 polypeptides/active
d APO4 signal transducer molecules that specifically interact
lamic domain of APO4 and detecting a change in level of APO4
e method is performed in vivo or in vitro. APO polypeptides
ul as immunogens for preparing antibodies. APO4 is also
agnosis/treatment of developmental or gestational
s. APO8 was transfected to human breast carcinoma cell line
duced apoptosis

AA;

16.2%; Score 46; DB 2; Length 211;
larity 100.0%; Pred. No. 9.6e-34;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LRLCQVSGLLALRPGSSLRITLTPWAHLKAAPFLTYGLFQVH 284
|||||
LRLCQVSGLLALRPGSSLRITLTPWAHLKAAPFLTYGLFQVH 211

adard; protein; 225 AA.

(first entry)

tumour necrosis factor related ligand (TRELL).

r necrosis factor related ligand; tnfr; treatment; cancer;
isease; immune system; stimulation; suppression;
ion.

Location/Qualifiers
1. 21

/note= "hydrophobic, transmembrane domain"

97WO-US013945.

96US-0023541P.

96US-0028515P.

97US-0040820P.

EN INC.
GENEVA FACULTY MEDICINE.

PI Chicheportiche Y, Browning JL;
XX WPI; 1998-145619/13.
DR N-PSDB; AAV18539.
XX
PT Tumour necrosis factor related ligand - useful for, e.g. treatir
PT auto-immune disease and immune responses to tissue grafts.
XX
XX Claim 12; Page 48-50; 69pp; English.

XX The sequence is that of mouse tumour necrosis factor related lig
CC (TRELL). TRELL or active fragments can be included with a carrie
CC pharmaceutical compositions to treat cancer, autoimmune diseases
CC immune responses to tissue grafts, or to stimulate or suppress t
CC system. It is useful to screen for TRELL receptors, by labelling
CC detectable label and screening compositions for binding. Agents
CC interfering with TRELL-receptor binding can also be screened for
CC then be administered, optionally with interferon- gamma, to indu
CC death or treat, suppress or alter immune responses (especially i
CC human adenocarcinoma cells) involving a signal pathway between t
CC its receptor. It's coding sequence can be used in gene therapy f
CC related disorders in mammals (especially humans), e.g. tumours,
CC autoimmune and inflammatory diseases or inherited genetic disord
CC introducing into cells, and expressing, therapeutically effectiv
CC of a vector, e.g. a virus comprising a gene encoding TRELL. It m
CC be of use in the preparation of prepare probes for screening
CC natural/synthetic DNAs for TRELL-encoding sequences and for anti
CC therapy
XX
XX SQ Sequence 225 AA;

Query Match 11.3%; Score 32; DB 2; Length 225;
Best Local Similarity 100.0%; Pred. No. 7.6e-21;
Matches 32; Conservative 0; Mismatches 0; Indels 0;

QY 139 RRAIAAHYEVHPRPGDGAQAGVDGTVSGWEE 170
|||||
Db 80 RRAIAAHYEVHPRPGDGAQAGVDGTVSGWEE 111

RESULT 17

AAB07527

ID AAB07527 standard; protein; 225 AA.

XX AAB07527;

XX 20-OCT-2000 (first entry)

XX Amino acid sequence of a soluble recombinant murine TWEAK protei
DE TWEAK protein; immunological disorder; immune response; inflama
XX TWEAK blocking agent; autoimmune disease; organ transplant rejec
KW Graft-versus-Host disease; GVHD; lymphoid cell malignancy; shock
XX
XX Mus sp.

XX WO200042073-A1.

XX 20-JUL-2000.

XX 14-JAN-2000; 2000WO-US001044.

XX 15-JAN-1999; 99US-0116168P.

XX (BIOJ) BIOGEN INC.

XX Rennert P;

XX WPI; 2000-476036/41.

XX Preventing and treating immune responses using modulators, espec:
PT antibodies, of TWEAK, TWEAK receptors and TWEAK ligands, useful
PT treating e.g. inflammation and graft versus host disease.

ig 1; 45pp; English.

sequence represents a TWEAK protein. The specification method for preventing or treating an immunological disorder using an immune response in an animal. The method comprises a TWEAK blocking agent. The method may be used for treating immune disorders associated with inappropriate d/or activity of TWEAK. These disorders include autoimmune and chronic inflammation, organ transplant rejection, Host disease (GVHD), lymphoid cell malignancies, septic and shock, loss of immune responsiveness (as seen in human HIV virus (HIV) infections) and failure of the immune response growth

AA;

11.3%; Score 32; DB 3; Length 225;
arity 100.0%; Pred. No. 7.6e-21;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

AAHYEHVPRPGQGAQAGVDGTVSGWEE 170
|||||
AAHYEHVPRPGQGAQAGVDGTVSGWEE 111

hard; protein; 249 AA.

(first entry)

AK.

EAK; TNF relatedness and weak ability to induce cell death;
scrosis factor; TWEAK; fibrosis; cardiac disease;
lung disease; kidney disease; skin disease;
le disease; adipose tissue disease;
nal tract disease; pancreatic disease;
organ disease; neural disease; cartilage disease;
connective tissue disease; cellular death; hepatotropic;
l; gastrointestinal; osteopathic.

12.

003WO-US011350.

002US-037161P.

INC.

ubowski A, Zheng T, Hahn K;

56/78.

13.

EAK-related condition, e.g. liver, gastrointestinal, kidney,
ic, cartilage or neural tissue condition in a subject
nistering to the subject a TWEAK agonist or antagonist.

ID NO 1; 120pp; English.

sequence is murine transmembrane FL-TWEAK (TNF relatedness
ty to induce cell death, where TNF is Tumour Necrosis
is a member of the TNF family. TWEAK agonists or
e useful for treating a TWEAK-related condition, e.g.
liac disease; liver disease; lung disease; kidney disease;

CC skin disease; skeletal muscle disease; adipose tissue disease;
CC gastrointestinal tract disease; pancreatic disease; reproductive
CC disease; neural disease; cartilage disease; bone disease; connect
CC tissue disease; cellular death; and a pathological condition of
CC expressing a TWEAK receptor.

SQ Sequence 249 AA;

Query Match 11.3%; Score 32; DB 7; Length 249;
Best Local Similarity 100.0%; Pred. No. 8.4e-21;
Matches 32; Conservative 0; Mismatches 0; Indels 0; G

Qy 139 RRAIAHYEHVPRPGQGAQAGVDGTVSGWEE 170
|||||
Db 104 RRAIAHYEHVPRPGQGAQAGVDGTVSGWEE 135

RESULT 19

AAG01265

ID AAG01265 standard; protein; 58 AA.

AC AAG01265;

DT 06-OCT-2000 (first entry)

DE Human secreted protein, SEQ ID NO: 5346.

KW Human; 5' EST; expressed sequence tag; secreted protein; cDNA iso
KW gene therapy; chromosome mapping.

OS Homo sapiens.

XX EP1033401-A2.

XX 06-SEP-2000.

XX 21-FEB-2000; 2000EP-00200610.

XX 26-FEB-1999; 99US-0122487P.

XX (GEST) GENSET.

XX Dumas Milne Edwards J, Duclert A, Giordano J;

XX WPI: 2000-50038-/45.

XX N-PSDB; AAC01272.

PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for
PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and fo
PT diagnostic, forensic, gene therapy and chromosome mapping procedu

XX Claim 13; SEQ ID NO 5346; 71pp + Sequence Listing; English.

CC The present sequence is a polypeptide encoded by one of a large n
CC 5' ESTs derived from mRNAs encoding secreted proteins. The 5' EST
CC prepared from total human RNAs or polyA+ RNAs derived from 30 dif
CC tissues. EST sequences usually correspond mainly to the 3' untran
CC region (UTR) of the mRNA because they are often obtained from Oli
CC primed cDNA libraries. Such ESTs are not well suited for isolatin
CC sequences derived from the 5' ends of mRNAs and even in those cas
CC longer cDNA sequences have been obtained, the full 5' UTR is rare
CC included. 5' ESTs are derived from mRNAs with intact 5' ends and
CC therefore be used to obtain full length cDNAs and genomic DNAs. 5'
CC are also used in diagnostic, forensic, gene therapy and chromosom
CC mapping procedures. They are used to obtain upstream regulatory s
CC and to design expression and secretion vectors

XX Sequence 58 AA;

Query Match

Best Local Similarity 3.2%; Score 9; DB 3; Length 58;

Matches 9; Conservative 0; Mismatches 0; Indels 0; G

9
5

standard; protein; 58 AA.

(first entry)

ed protein, SEQ ID NO: 5347.

†; expressed sequence tag; secreted protein; cDNA isolation; chromosome mapping.

2000EP-00200610.

99US-0122487P.

1.

Edwards J, Duclert A, Giordano J;

0381/45.

272.

acid that is a 5' expressed sequence tag (5' EST) for cDNAs and genomic DNAs that correspond to 5'ESTs and for forensic, gene therapy and chromosome mapping procedures.

ID NO 5347; 71pp + Sequence Listing; English.

sequence is a polypeptide encoded by one of a large number of mRNAs encoding secreted proteins. The 5' ESTs were a total human RNAs or polyA+ RNAs derived from 30 different sequences usually correspond mainly to the 3' untranslated sequences of the mRNA because they are often obtained from oligo-dT libraries. Such ESTs are not well suited for isolating cDNA derived from the 5' ends of mRNAs and even in those cases where sequences have been obtained, the full 5' UTR is rarely obtained. ESTs are derived from mRNAs with intact 5' ends and can be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are used in diagnostic, forensic, gene therapy and chromosome mapping studies. They are used to obtain upstream regulatory sequences and expression and secretion vectors.

A:

3.28; Score 9; DB 3; Length 58;

urity 100.0%; Pred. No. 2.9;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LD FEIS 9 54

Standard: protein: 365 AA.

(first entry)

XX	Human D1THP receptor.
XX	
DE	Human; dithp; diagnostic and therapeutic polynucleotide; diagnos
DE	cancer; cell proliferative disorder; autoimmune disorder;
KW	cancer; cell proliferative disorder; autoimmune disorder;
KW	inflammatory disorder; infection; hormonal disorder; metabolic d
KW	neurological disorder; gastrointestinal disorder; transport diso
KW	connective tissue disorder; drug screening; proteome analysis;
KW	gene therapy; antisense therapy; genotyping; transgenic animal;
KW	disease model; toxicological testing; transcript imaging; recept
XX	
OS	Homo sapiens.
XX	
PN	WO200297031-A2.
XX	
PD	05-DEC-2002.
XX	
PF	27-MAR-2002; 2002WO-US010056.
XX	
PR	28-MAR-2001; 2001US-0279619P.
PR	29-MAR-2001; 2001US-0280067P.
PR	29-MAR-2001; 2001US-0280068P.
PR	16-MAY-2001; 2001US-0291280P.
PR	17-MAY-2001; 2001US-0291829P.
PR	17-MAY-2001; 2001US-0291849P.
PR	19-JUN-2001; 2001US-0299428P.
PR	20-JUN-2001; 2001US-0299776P.
PR	20-JUN-2001; 2001US-0300001P.
XX	
PA	(INCY-) INCYTE GENOMICS INC.
XX	
PI	Daffo A, Jones AL, Tran AB, Dahl CR, Gietzen D, Chinn J;
PI	Duffour GE, Hillman JL, Yu JY, Tuason O, Yap PE, Amshay SR;
PI	Daugherty SC, Dam TC, Liu TF, Nguyen DA, Kleefeld V, Gersti
PI	Peralta CH, David MH, Lewis SA, Chen AJ, Panzer SR, Harris
PI	Flores V, Marwaha R, Lo A, Lan RY, Urashka ME;
PI	
DR	WPI; 2003-129518/12.
DR	N-PSDB; ACC46177.
XX	
PT	Novel human diagnostic and therapeutic polypeptide useful for id
PT	test compound which specifically binds to a polypeptide encoded i
PT	diagnostic and therapeutic polynucleotide, and to induce antibod
XX	
PS	Claim 27; SEQ ID NO 770; 591pp; English.
XX	
CC	The invention relates to novel human diagnostic and therapeutic
CC	polynucleotides designated dithp (ACC46080-ACC46749) and to thei
CC	proteins (D1THP; ABR41136-ABR41812). The invention also relates i
CC	polynucleotide sequences at least 90% identical to the dithp cDN
CC	sequences of the invention; recombinant vectors, host cells and
CC	transgenic organisms comprising a dithp nucleic acid sequence; t
CC	recombinant production of D1THP proteins; antibodies specific fo
CC	proteins; microarrays comprising dithp nucleic acid sequences; m
CC	detecting dithp nucleotide and protein sequences; methods of scr
CC	for compounds which specifically bind a D1THP protein; and metho
CC	assessing the toxicity of test compounds using a dithp hybridisa
CC	probe. Dithp nucleic acid sequences and D1THP proteins may be use
CC	diagnosis of a wide variety of conditions including cancer and o
CC	proliferative disorders; autoimmune or inflammatory disorders; b
CC	viral, fungal or parasitic infections; hormonal disorders; metabo
CC	disorders; neurological disorders; gastrointestinal disorders; t
CC	disorders; and connective tissue disorders. They may also be use
CC	screen for modulators of protein activity or gene expression. D1
CC	proteins can additionally be used in analysis of the proteome of
CC	or cell type and to induce antibodies. The dithp nucleic acids a
CC	additionally useful in somatic or germline gene therapy of the d
CC	mentioned above, as a source of antisense sequences, as a source
CC	probes and primers, in genotyping and identification of individu
CC	the generation of transgenic animal models of human disease or k
CC	humanised animals, in toxicological testing and in transcript i
CC	The present sequence represents a D1THP protein which has recept
CC	activity. Note: The sequence data for this patent did not form p
CC	

pecification, but was obtained in electronic format directly
ftp.wipo.int/pub/published_pct_sequences

AA;

3.2%; Score 9; DB 6; Length 365;
arity 100.0%; Pred. No. 17;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

GRGCE 50

|||||

GRGCE 30

dard; protein; 748 AA.

(first entry)

o acid sequence for GVs-9.

vaccae protein; antigen; T cell activation; cytokine;
1 maturation; infectious disease; immune disorder; cancer;
ystem; mycobacterial infection; allergy; tuberculosis;
oidosis; lung cancer; asthma; skin disorder; psoriasis;
czema; alopecia areata; skin cancer; basal carcinoma;
carcinoma; melanoma.

vaccae.

98WO-NZ000189.

97US-00996624.

97US-0097080.

97US-00997362.

98US-00095855.

98US-00156181.

98US-00205436.

IS RES & DEV CORP LTD.

n J, Visser ES, Skinner MA, Prestidge RL;

163/36.

368.

une response to an antigen.

209-210; 243pp; English.

provides heat-killed Mycobacterium vaccae, or recombinant
teins. The M. vaccae proteins may be employed to activate T
ural killer cells, to stimulate the production of cytokines,
e expression of co-stimulatory molecules on dendritic cells
, and to enhance dendritic cell maturation and function. The
be expressed by standard recombinant methodology.

1 compositions comprising the proteins or nucleic acid
oding the proteins can be used for the treatment,
nd detection of disorders including infectious diseases,
ers and cancer. In particular, the compounds and methods are
tment of diseases of the respiratory system, such as
infections, asthma, allergies, tuberculosis, leprosy,
nd lung cancers, and disorders of the skin such as
opic dermatitis, eczema, allergic contact dermatitis,
ta, and skin cancers such as basal carcinoma, squamous cell
melanoma

XX SQ Sequence 748 AA;

Query Match 3.2%; Score 9; DB 2; Length 748;
Best Local Similarity 100.0%; Pred. No. 34;
Matches 9; Conservative 0; Mismatches 0; Indels 0;

QY 60 ALGGLALA 68

|||||

Db 282 ALGGLALA 290

RESULT 23

ABB73512

ID ABB73512 standard; protein; 749 AA.

XX

AC ABB73512;

XX

DT 08-APR-2002 (first entry)

XX

DE M vaccae GVs-9 protein SEQ ID NO: 154.

XX

KW Skin disorder; psoriasis; atopic dermatitis; allergic contact der;
KW alopecia areata; skin cancer; Mycobacterium vaccae; melanoma; cyt
KW antipruritic; dermatological; antiinflammatory; antiallergic;
KW Th2 immune response; immunomodulatory.

XX

OS Mycobacterium vaccae.

XX

FN US6328978-B1.

XX

PD 11-DEC-2001.

XX

PF 02-JUN-1999; 99US-00324542.

XX

PR 23-DEC-1997; 97US-00997080.

XX

PA (GENE-) GENESIS RES & DEV CORP LTD.

XX

PI Watson JD, Tan PLJ, Prestidge R;

XX

DR WPI; 2002-138361/18.

XX

DR N-PSDE; ABL36274.

XX

Inhibiting skin inflammation associated with skin disorder e.g.
psoriasis, by administering composition comprising delipidated ar
deglycolipidated Mycobacterium vaccae cells or Mycobacterium vacc
culture filtrate.

XX

PS Example 6; Col 161-164; 116pp; English.

XX

The present invention relates to a method of inhibiting skin infl
associated with a skin disorder selected from psoriasis, atopic
dermatitis and allergic contact dermatitis, which involves admini
a composition containing delipidated and deglycolipidated Mycobac
vaccae cells or M. vaccae culture filtrate. The skin disorder to
CC treated may also include alopecia areata, and skin cancers such a
CC cell carcinoma, squamous cell carcinoma and melanoma. The composi
CC acts by inhibiting the Th2 immune response. The present sequence
CC protein described in the exemplification of the invention

XX

SQ Sequence 749 AA;

Query Match 3.2%; Score 9; DB 5; Length 749;

Best Local Similarity 100.0%; Pred. No. 34;
Matches 9; Conservative 0; Mismatches 0; Indels 0; G

QY 60 ALGGLALA 68

|||||

Db 282 ALGGLALA 290

RESULT 24

standard; protein; 54 AA.

(first entry)

erium acnes immunogenic protein #12759.

me; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
ophthalmitis; bone; joint; central nervous system; ELISA;
lesion; acne vulgaris; enzyme linked immunosorbent assay;
al; osteopathic; neuroprotectant.

erium acnes.

A2.

2001WO-US012865.

2000US-0199047P.

2000US-0208841P.

2000US-0216747P.

XA CORP.

Persing DH, Mitcham JL, Wang SS, Bhatia A;
e J, Zhang Y, Jen S, Carter D;

6774/71.

9552.

erium acnes polypeptides and nucleic acids useful for
against and diagnosing infections, especially useful for
e vulgaris.

EQ ID NO 13058; 1069pp; English.

U39105-AA068017 represent Propionibacterium acnes immunogenic
t. The proteins and their associated DNA sequences are used in
t, prevention and diagnosis of medical conditions caused by
e disorders include SAPHO syndrome (synovitis, acne,
hypertosis and osteomyelitis), uveitis and endophthalmitis.
also involved in infections of bone, joints and the central
em, however it is particularly involved in the inflammatory
ciated with acne vulgaris. A method for detecting the
absence of P. acnes in a patient comprises contacting a
binding agent that binds to the proteins of the invention
ing the amount of bound protein in the sample. The
may be used as antigens in the production of antibodies
P. acnes proteins. These antibodies can be used to
expression and activity of P. acnes polypeptides and
eat P. acnes infections. The antibodies may also be used as
gents for determining P. acnes presence, for example, by
d immunosorbent assay (ELISA). Note: The sequence data for
did not form part of the printed specification, but was
electronic format directly from WIPO at
/pub/published_pct_sequences

AA;

larity 2.8%; Score 8; DB 4; Length 54;
100.0%; Pred. No. 23;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

PLPRS 18

|||||

PLPRS 28

ABM48382

ID ABM48382 standard; protein; 54 AA.

XX AC ABM48382;

XX DT 20-OCT-2003 (first entry)

XX DE Propionibacterium acnes predicted ORF-encoded polypeptide #13058

XX KW Acne vulgaris; antiseborrheic; dermatological; antibacterial;

XX KW Immunostimulant; immune response; vaccine.

XX OS Propionibacterium acnes.

XX PN WO2003033515-A1.

XX PD 24-APR-2003.

XX PF 11-OCT-2002; 2002WO-US032727.

XX PR 15-OCT-2001; 2001US-00978825.

XX PA (CORI-) CORIXA CORP.

XX PI Mitcham JL, Skeiky YAW, Persing DH, Bhatia A, Maisonneuve JI

PI Zhang Y, Wang S, Jen S, Lodes MJ, Benson DR, Jones R, Cart

PI Barth B, Vallieve-Douglass J;

XX WPI; 2003-391799/36.

DR N-PSDB; ACF64481.

XX New Propionibacterium acnes polypeptides and polynucleotides enc

PT polypeptide, useful for diagnosing, preventing or treating acne

PT or for stimulating an immune response specific for a P. acnes pr

XX Example 1; SEQ ID NO 13058; 1481pp; English.

XX The invention relates to an isolated polynucleotide (ACF64435-AC

CC encoding a Propionibacterium acnes protein. The invention also r

CC polypeptides encoded by the polynucleotides (ABM35624-ABM4536)

CC immunogenic fragments of P. acnes polypeptides. The invention

CC additionally encompasses expression vectors and host cells compr

CC polynucleotide of the invention; antibodies against polypeptides

CC invention; fusion proteins comprising a polypeptide of the inven

CC method for stimulating an immune response specific for a P. acne

CC polypeptide and an isolated T cell population comprising T cells

CC via this method; a vaccine composition (comprising P. acnes poly

CC polynucleotides, antibodies, fusion proteins, T cell populations

CC antigen-presenting cells that express the polypeptide); a method

CC for detecting or determining the presence or absence of P. acnes

CC patient; and a method for inhibiting the development of P. acnes

CC proteins. The P. acnes polypeptides, polynucleotides, antibodies,

CC proteins, T cell populations or antigen-presenting cells that ex

CC polypeptides are useful for diagnosing, preventing or treating a

CC vulgaris, or for stimulating an immune response specific for a P

CC protein. The polynucleotides can also be used as probes or prime

CC nucleic acid hybridisation. The vaccine composition is useful fo

CC stimulation of an immune response against P. acnes, or for treat

CC and the kit is useful for performing a diagnostic assay. The pre

CC sequence represents a polypeptide predicted to be encoded by an

CC reading frame) contained within the P. acnes polynucleotides of an

CC invention. Note: The sequence data for this patent did not form

CC the printed specification, but was obtained in electronic format

CC from WIPO at ftp.wipo.int/pub/published_pct_sequences

XX Sequence 54 AA;

XX Query Match 2.8%; Score 8; DB 6; Length 54;

Best Local Similarity 100.0%; Pred. No. 23;

Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 11 RRLPLPRS 18

|||||

Db

LPRS 28

DB

XX

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XX

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XX

dard; protein; 55 AA.

(first entry)

encoded by probe for measuring cervical gene expression.

microarray; gene expression; cervical epithelial cell;

er.

2.

2001WO-US000670.

2000US-0180312P.

2000US-0207456P.

2000US-00608408.

2000US-00632366.

2000US-0234687P.

2000US-0236359P.

2000GB-00024263.

ULAR DYNAMICS INC.

zel DK, Chen W, Rank DR;

901/53.

derived single exon nucleic acid probes useful for analyzing
on in human cervical epithelial cells.

ID NO 25684; 487pp; English.

vention relates to human single exon nucleic acid probes
110068-AA128459). The present sequence is a peptide encoded
robe. The SNPs are derived from human HeLa cells. The SNPs
o produce a single exon microarray, which can be used for
an gene expression in a sample derived from human cervical
lles. By measuring gene expression, the probes are therefore
ding and/or staging of diseases of the cervix, notably
er. Note: The sequence data for this patent did not form
rinted specification, but was obtained in electronic format
WIPO at ftp.wipo.int/pub/published_pct_sequences

a;

arity 2.8%; Score 8; DB 4; Length 55;

nservative 100.0%; Pred.No. 23;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

LALA 68

LALA 19

dard; peptide; 55 AA.

(first entry)

DE Peptide #10143 encoded by human foetal liver single exon probe.

XX Human; foetal liver; gene expression; single exon nucleic acid p;

XX Homo sapiens.

OS WO200157277-A2.

XX 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US000669.

XX 04-FEB-2000; 2000US-0180312P.

XX 26-MAY-2000; 2000US-0207456P.

XX 30-JUN-2000; 2000US-00608408.

XX 03-AUG-2000; 2000US-00632366.

XX 21-SEP-2000; 2000US-0234687P.

XX 27-SEP-2000; 2000US-0236359P.

XX 04-OCT-2000; 2000GB-00024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-483447/52.

XX Human genome-derived single exon nucleic acid probes useful for e

XX gene expression in human fetal liver.

XX Claim 27; SEQ ID NO 35272; 639pp + Sequence Listing; English.

XX The invention relates to a single exon nucleic acid probe for mea

XX human gene expression in a sample derived from human foetal live;

XX single exon nucleic acid probes may be used for predicting, measu

XX displaying gene expression in samples derived from human fetal li

XX present sequence is a peptide encoded by a single exon nucleic ac

XX of the invention. Note: The sequence data for this patent did not

XX part of the printed specification, but was obtained in electronic

XX directly from WIPO at ftp.wipo.int/pub/published_pct_sequences

XX Sequence 55 AA;

XX Query Match 2.8%; Score 8; DB 4; Length 55;

XX Best Local Similarity 100.0%; Pred.No. 23;

XX Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 61 LGUGLALA 68

Db 12 LGUGLALA 19

RESULT 28

AAM36451

ID AAM36451 standard; protein; 55 AA.

XX AAM36451;

XX 17-OCT-2001 (first entry)

XX Peptide #10488 encoded by probe for measuring placental gene expr

XX Probe; microarray; human; placenta; antenatal diagnosis;

XX genetic disorder.

XX Homo sapiens.

XX WO200157272-A2.

XX 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US000663.

XX 04-FEB-2000; 2000US-0180312P.

XX

2000US-0207456P.
2000US-00608408.
2000US-00632366.
2000US-0234687P.
2000US-0236359P.
2000GB-00024263.

MOLECULAR DYNAMICS INC.

nzal DK, Chen W, Rank DR;

8897/53.

--derived single exon nucleic acid probes useful for analyzing
ion in human placenta.

Q ID NO 36720; 654pp; English.

invention relates to single exon nucleic acid probes (SENP:
-AA157546). The present sequence is a peptide encoded by one
The probes are useful for producing a microarray for
measuring and displaying gene expression in samples derived
ilacenta. The probes are useful for antenatal diagnosis of
c disorders

AA;

larity 2.8%; Score 8; DB 4; Length 55;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GLALA 68

GLALA 19

ndard; protein; 55 AA.

(first entry)

2 encoded by probe for measuring heart cell gene expression.

expression; heart; microarray; vascular system;
ar disease; hypertension; cardiac arrhythmia;
eart disease.

A2.

2001WO-US000666.

2000US-0180312P.
2000US-0207456P.
2000US-00608408.
2000US-00632366.
2000US-0234687P.
2000US-0236359P.
2000GB-00024263.

CULAR DYNAMICS INC.

nzal DK, Chen W, Rank DR;

8899/53.

nucleic acid probes for analyzing gene expression in human

PT hearts.

XX Claim 15; SEQ ID NO 27753; 530pp; English.

XX The present invention relates to single exon nucleic acid probes
XX measuring human gene expression in a sample derived from human
XX ABA21535-ABA41305). The present sequence is a protein encoded by
XX probe. The probes may be used for predicting, measuring and dis
XX gene expression in samples derived from the human heart via mic
XX By measuring gene expression, the probes are useful for predict
XX diagnosing, grading, staging, monitoring and prognosing disease
XX human heart and vascular system e.g. cardiovascular disease,
XX hypertension, cardiac arrhythmias and congenital heart disease.
XX sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly f
XX at ftp.wipo.int/pub/published_pct_sequences

SQ Sequence 55 AA;

Query Match 2.8%; Score 8; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

Qy 61 LGGLGALA 68

Db 12 LGGLGALA 19

RESULT 30

AAM76342

ID AAM76342 standard; protein; 55 AA.

XX AAM76342;

DT 06-NOV-2001 (first entry)

XX Human bone marrow expressed probe encoded protein SEQ ID NO: 366
XX Human; bone marrow expressed exon; gene expression analysis; pr
XX microarray; cancer; leukaemia; lymphoma; myeloma.

OS Homo sapiens.

XX WO200157276-A2.

XX 09-AUG-2001.

PD 30-JAN-2001; 2001WO-US000668.

XX 04-FEB-2000; 2000US-0180312P.

PR 26-MAY-2000; 2000US-0207456P.

PR 30-JUN-2000; 2000US-00608408.

PR 03-AUG-2000; 2000US-00632366.

PR 21-SEP-2000; 2000US-0234687P.

PR 27-SEP-2000; 2000US-0236359P.

PR 04-OCT-2000; 2000GB-00024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-488900/53.

XX Human genome-derived single exon nucleic acid probes useful for
XX gene expression in human bone marrow.

XX Example 4; SEQ ID NO 36648; 658pp + Sequence Listing; English.

XX The present invention provides a number of single exon nucleic a
XX probes which are derived from genomic sequences expressed in
XX bone marrow. They can be used to measure gene expression in bone
XX samples, which may enable the improved diagnosis and treatment o
XX such as lymphoma, leukaemia and myeloma. The present sequence is

ad by one of the probes of the invention

2.8%; Score 8; DB 4; Length 55;
arity 100.0%; Pred. No. 23;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
ALA 68
|||||
ALA 19

iard; protein; 55 AA.

(first entry)

pressed single exon probe encoded protein SEQ ID NO: 35633.
expressed exon; gene expression analysis; probe; microarray;
isease; multiple sclerosis; schizophrenia; epilepsy; cancer.

2.

2001WO-US000667.

2000US-0180312P.
2000US-0207456P.
2000US-00608408.
2000US-00632366.
2000US-0234687P.
2000US-0236359P.
2000GB-00024263.

ULAR DYNAMICS INC.

zel DK, Chen W, Rank DR;

446/52.

ucleic acid probes for analyzing gene expression in human

Q ID NO 35633; 650pp + Sequence Listing; English.

vention provides a number of single exon nucleic acid
are derived from genomic sequences expressed in the human
an be used to measure gene expression in brain cell samples,
ble the diagnosis and improved treatment of nervous system
as Alzheimer's disease, multiple sclerosis, schizophrenia, f
cancers. The present sequence is a protein encoded by one of
the invention

2.8%; Score 8; DB 4; Length 55;
arity 100.0%; Pred. No. 23;
onservative 0; Mismatches 0; Indels 0; Gaps 0;
ALA 68
|||||
ALA 19

ABG58050
ID ABG58050 standard; peptide; 55 AA.
XX
AC ABG58050;
XX
DT 25-FEB-2003 (first entry)
XX
DE Human liver peptide, SEQ ID No 36698.
XX
XX Human; liver; cirrhosis; hyperlipoproteinaemia; hyperlipidaemia;
XX hypercholesterolaemia; coronary heart disease.
XX
OS Homo sapiens.
XX
PN WO200157273-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US000664.
XX
PR 04-FEB-2000; 2000US-0180312P.
PR 26-MAY-2000; 2000US-0207456P.
PR 30-JUN-2000; 2000US-00608408.
PR 03-AUG-2000; 2000US-00632366.
PR 21-SEP-2000; 2000US-0234687P.
PR 27-SEP-2000; 2000US-0236359P.
PR 04-OCT-2000; 2000GB-00024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI; 2001-488898/53.
XX
XX Human genome-derived single exon nucleic acid probes useful for a
gene expression in human adult liver.
XX
XX Claim 27; SEQ ID NO 36698; 659pp; English.
XX
XX The invention relates to a single exon nucleic acid probe (SENP)
measuring human gene expression in a sample derived from human ac
liver, comprising one of 13109 defined nucleotide sequences giver
specification (or complements/ fragments). The probe hybridises a
stringency to a nucleic acid molecule expressed in the human adul
(I) may be used for predicting, measuring and displaying gene exp
in samples derived from human adult liver. The genes identified
involved in genetic liver diseases such as cirrhosis,
hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia
associated with coronary heart disease. ABG47348-ABG59930 repres
liver single exon encoded peptides of the invention. Note: The se
information for this patent does not appear in the printed speci
but was obtained in electronic format directly from WIPO at
ftp.wipo.int/pub/published_pct_sequences
XX
XX Sequence 55 AA;
XX
XX Query Match 2.8%; Score 8; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 8; Conservative 0; Mismatches 0; Indels 0;
QY 61 LGLGLALA 68
|||||
DB 12 LGLGLALA 19
RESULT 33
ID ABG45635
XX
XX AC ABG45635;
XX
XX DT 19-AUG-2002 (first entry)
XX

e encoded by genome-derived single exon probe SEQ ID 35300.
 e exon probe; asthma; lung cancer; COPD; ILD;
 ractive pulmonary disease; interstitial lung disease;
 opathic pulmonary fibrosis; neurofibromatosis;
 erosis; Gaucher's disease; Niemann-Pick disease;
 dlak syndrome; sarcoidosis; pulmonary haemosiderosis;
 sticyctosis; lymphangioleiomyomatosis; Karagener syndrome;
 veolar proteinosis; fibrocystic pulmonary dysplasia;
 ary dyskinesia; pulmonary hypertension;
 rane disease.

A2.

2001WO-US000665.

2000US-0180312P.
 2000US-0207456P.
 2000US-00608408.
 2000US-00632366.
 2000US-0234687P.
 2000US-0236359P.
 2000GB-00024263.

TULAR DYNAMICS INC.

zel DK, Chen W, Rank DR;

1183/15.

irreversible set of single exon nucleic acid probes, used to
 expression in human lung samples.

2 ID NO 35300; 634pp; English.

1 relates to a spatially-addressable set of single exon
 probes for measuring gene expression in a sample deriv
 ing comprising single exon nucleic acid probes having one of
 ; acid sequences mentioned in the specification, or their
 or the 12387 open reading frames derived from the 12614
 included are a microarray comprising the novel set of probes
 set of probes which hybridise at high stringency to a nucleic
 id in the human lung; measuring gene expression in a sample
 human lung, comprising (a) contacting the array with a
 ; detectably labeled nucleic acids derived from human lung
 measuring the label detectably bound to each probe of the
 .fying exons in a eukaryotic genome, comprising (a)
 ly predicting at least one exon from genomic sequences of
 ; and (b) detecting specific hybridisation of detectably
 ic acids from eukaryotic lung mRNA, to a single exon probe,
 ment identical to the predicted exon, the probe is included
 mentioned microarray; assigning exons to a single gene,
 u) identifying exons from genomic sequence by the method
 measuring the expression of each of the exons in several
 or cell types using hybridisation to a single exon
 having a probe with the exon, where a common pattern of
 ; the exons in the tissues and/or cell types indicates that
 uld be assigned to a single gene; a peptide comprising one
 ences, mentioned in the specification, or encoded by the
 reading frames (ORF). The probes are used for gene expression
 l for identifying exons in a gene, particularly using human
 mRNA and for the study of lung diseases such as asthma, lung
 ic obstructive pulmonary disease (COPD), interstitial lung
 , familial idiopathic pulmonary fibrosis, neurofibromatosis,
 ;rosis, Gaucher's disease, Niemann-Pick disease, Hermansky-
 me, sarcoidosis, pulmonary haemosiderosis, pulmonary
 , lymphangioleiomyomatosis, pulmonary alveolar proteinosis,
 idrome, fibrocystic pulmonary dysplasia, primary ciliary
 pulmonary hypertension and hyaline membrane disease. The

CC present sequence is a peptide/protein encoded by a single exon I
 CC the invention. Note: The sequence data for this patent did not i
 CC of the printed specification, but was obtained in electronic for
 CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 55 AA;

Query Match 2.8%; Score 8; DB 5; Length 55;
 Best Local Similarity 100.0%; Pred. No. 23;
 Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 61 LGLGLALA 68
 |||||
 Db 12 LGLGLALA 19

RESULT 34

AAW21621
 ID AAW21621 standard; protein; 65 AA.

XX

AC AAW21621;

XX 12-OCT-2001 (first entry)

DT

XX Peptide #8055 encoded by probe for measuring cervical gene expre
 DE Probe; human; microarray; gene expression; cervical epithelial c
 XX cervical cancer.
 XX Homo sapiens.

OS

XX WO200157278-A2.

PN

XX 09-AUG-2001.

PD

PF 30-JAN-2001; 2001WO-US000670.

XX

PR 04-FEB-2000; 2000US-0180312P.

PR 26-MAY-2000; 2000US-0207456P.

PR 30-JUN-2000; 2000US-00608408.

PR 03-AUG-2000; 2000US-00632366.

PR 21-SEP-2000; 2000US-0234687P.

PR 27-SEP-2000; 2000US-0236359P.

PR 04-OCT-2000; 2000GB-00024263.

XX

PA (MOLE-) MOLECULAR DYNAMICS INC.

XX

PI Penn SG, Hanzel DK, Chen W, Rank DR;

XX

XX WPI; 2001-488901/53.

DR

PT Human genome-derived single exon nucleic acid probes useful for
 PT gene expression in human cervical epithelial cells.

XX

PS Claim 27; SEQ ID NO 26447; 487pp; English.

XX

CC The present invention relates to human single exon nucleic acid I
 CC (SENP; see AAI10068-AA128459). The present sequence is a peptide
 CC by one such probe. The SENPs are derived from human HeLa cells.
 CC can be used to produce a single exon microarray, which can be use
 CC measuring human gene expression in a sample derived from human ce
 CC epithelial cells. By measuring gene expression, the probes are t
 CC useful in grading and/or staging of diseases of the cervix, notat
 CC cervical cancer. Note: The sequence data for this patent did not
 CC part of the printed specification, but was obtained in electronic
 CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
 XX

SQ Sequence 65 AA;

Query Match

Best Local Similarity 2.8%; Score 8; DB 4; Length 65;

Matches 8; Conservative 0; Mismatches 0; Indels 0;

LALA 68
||||
LALA 29

lard; peptide; 65 AA.

(first entry)

7 encoded by human foetal liver single exon probe.

liver; gene expression; single exon nucleic acid probe.

2.

2001WO-US000669.

2000US-0180312P.
2000US-0207456P.
2000US-00608408.
2000US-00632366.
2000US-0234687P.
2000US-0236359P.
2000GB-00024263.

JLAR DYNAMICS INC.

zel DK, Chen W, Rank DR;

147/52.

derived single exon nucleic acid probes useful for analyzing
in human fetal liver.

ID NO 36616; 639pp + Sequence Listing; English.

relates to a single exon nucleic acid probe for measuring
expression in a sample derived from human foetal liver. The
nucleic acid probes may be used for predicting, measuring and
expression in samples derived from human fetal liver. The
probe is a peptide encoded by a single exon nucleic acid probe
on. Note: The sequence data for this patent did not form
part of the specification, but was obtained in electronic format
WIPO at ftp.wipo.int/pub/published_pct_sequences

2.8%; Score 8; DB 4; Length 65;

identity 100.0%; Pred. No. 27;

conservative 0; Mismatches 0; Indels 0; Gaps 0;

LALA 68
||||
LALA 29

lard; protein; 65 AA.

(first entry)

7 encoded by probe for measuring placental gene expression.

XX Probe; microarray; human; placenta; antenatal diagnosis;
KW genetic disorder.
XX
XX Homo sapiens.
OS
XX WO200157272-A2.
PN
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US000663.
PF
XX
XX 04-FEB-2000; 2000US-0180312P.
PR
XX 26-MAY-2000; 2000US-0207456P.
PR
XX 30-JUN-2000; 2000US-00608408.
PR
XX 03-AUG-2000; 2000US-00632366.
PR
XX 21-SEP-2000; 2000US-0234687P.
PR
XX 27-SEP-2000; 2000US-0236359P.
PR
XX 04-OCT-2000; 2000GB-00024263.
PR
XX (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX WPI; 2001-488897/53.
XX
DR Human genome-derived single exon nucleic acid probes useful for a
XX gene expression in human placenta.
PT
PT Claim 27; SEQ ID NO 38192; 654pp; English.
XX
XX The present invention relates to single exon nucleic acid probes
CC see AA131315-AA157546). The present sequence is a peptide encoded
CC such probe. The probes are useful for producing a microarray for
CC predicting, measuring and displaying gene expression in samples d
CC from human placenta. The probes are useful for antenatal diagnosi
CC human genetic disorders
XX
XX Sequence 65 AA;
SQ
Query Match 2.8%; Score 8; DB 4; Length 65;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G
QY 61 LGLGLALA 68
Db 22 LGLGLALA 29
|||||
|||||
RESULT 37
ABB26890
ID ABB26890 standard; protein; 65 AA.
XX
XX ABB26890;
AC
XX
XX 23-JAN-2002 (first entry)
DT
XX
XX Protein #889 encoded by probe for measuring heart cell gene expr
DE
XX
XX Human; gene expression; heart; microarray; vascular system;
KW cardiovascular disease; hypertension; cardiac arrhythmia;
KW congenital heart disease.
XX
XX Homo sapiens.
OS
XX WO200157274-A2.
PN
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US000666.
PF
XX
XX 04-FEB-2000; 2000US-0180312P.
PR
XX 26-MAY-2000; 2000US-0207456P.
PR

2000US-00608408.
2000US-00632366.
2000US-0234687P.
2000US-0236359P.
2000GB-00024263.

ULAR DYNAMICS INC.

azel DK, Chen W, Rank DR;

3899/53.

nucleic acid probes for analyzing gene expression in human

Q ID NO 28660; 530pp; English.

invention relates to single exon nucleic acid probes for
nan gene expression in a sample derived from human heart (see
11305). The present sequence is a protein encoded by one such
cubes may be used for predicting, measuring and displaying
ion in samples derived from the human heart via microarrays.
gene expression, the probes are useful for predicting, the
grading, staging, monitoring and prognosing diseases of the
and vascular system e.g. cardiovascular disease,
cardiac arrhythmias and congenital heart disease. Note: The
a for this patent did not form part of the printed
n, but was obtained in electronic format directly from WIPO
nt./pub/published_pct_sequences

LA;

2.8%; Score 8; DB 4; Length 65;
arity 100.0%; Pred. No. 27;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;

HLALA 68
|||||
HLALA 29

ard; protein; 65 AA.

(first entry)

arrow expressed probe encoded protein SEQ ID NO: 38012.

arrow expressed exon; gene expression analysis; probe;
ancer; leukaemia; lymphoma; myeloma.

2.

2001WO-US000668.

2000US-0180312P.
2000US-0207456P.
2000US-00608408.
2000US-00632366.
2000US-0234687P.
2000US-0236359P.
2000GB-00024263.

ULAR DYNAMICS INC.

azel DK, Chen W, Rank DR;

XX

DR WPI; 2001-488900/53.

XX

Human genome-derived single exon nucleic acid probes useful for
gene expression in human bone marrow.

XX

PS Example 4; SEQ ID NO 38012; 658pp + Sequence Listing; English.

XX

The present invention provides a number of single exon nucleic a
probes which are derived from genomic sequences expressed in the
bone marrow. They can be used to measure gene expression in bone
samples, which may enable the improved diagnosis and treatment o
such as lymphoma, leukaemia and myeloma. The present sequence is
protein encoded by one of the probes of the invention

XX

SQ Sequence 65 AA;

Query Match 2.8%; Score 8; DB 4; Length 65;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 61 LGLGLALA 68

Db 22 LGLGLALA 29

RESULT 39

AAM64984

ID AAM64984 standard; protein; 65 AA.

XX

AC AAM64984;

XX

DT 05-NOV-2001 (first entry)

XX

DE Human brain expressed single exon probe encoded protein SEQ ID N

XX

KW Human; brain expressed exon; gene expression analysis; probe; mi

KW

Alzheimer's disease; multiple sclerosis; schizophrenia; epilepsy

XX

OS Homo sapiens.

XX

PN WO200157275-A2.

XX

PD 09-AUG-2001.

XX

PF 30-JAN-2001; 2001WO-US000667.

XX

PR 04-FEB-2000; 2000US-0180312P.

PR

26-MAY-2000; 2000US-0207456P.

PR

30-JUN-2000; 2000US-00608408.

PR

03-AUG-2000; 2000US-00632366.

PR

21-SEP-2000; 2000US-0234687P.

PR

27-SEP-2000; 2000US-0236359P.

PR

04-OCT-2000; 2000GB-00024263.

XX

PA (MOLE-) MOLECULAR DYNAMICS INC.

XX

PI Penn SG, Hanzel DK, Chen W, Rank DR;

XX

DR WPI; 2001-483446/52.

XX

Single exon nucleic acid probes for analyzing gene expression in

PT

brains.

XX

Example 4; SEQ ID NO 37089; 650pp + Sequence Listing; English.

XX

The present invention provides a number of single exon nucleic ac
probes which are derived from genomic sequences expressed in the
brain. They can be used to measure gene expression in brain cell
which may enable the diagnosis and improved treatment of nervous
diseases such as Alzheimer's disease, multiple sclerosis, schizof
epilepsy and cancers. The present sequence is a protein encoded t
the probes of the invention

2.8%; Score 8; DB 4; Length 65;
 100.0%; Pred. No. 27;
 0; Mismatches 0; Indels 0; Gaps 0;

ALA 68
 ||||
 ALA 29

lard; peptide; 65 AA.

first entry)

ptide, SEQ ID No 38009.

cirrhosis; hyperlipoproteinaemia; hyperlipidaemia;
 olaemia; coronary heart disease.

001WO-US000664.

000US-0180312P.
 000US-0207456P.
 000US-00608408.
 000US-00632366.
 000US-0234587P.
 000US-0236359P.
 000GB-00024263.

LAR DYNAMICS INC.

el DK, Chen W, Rank DR;
 98/53.

erived single exon nucleic acid probes useful for analyzing
 n in human adult liver.

ID NO 38009; 658pp; English.

relates to a single exon nucleic acid probe (SENP) (I) for
 n gene expression in a sample derived from human adult
 ing one of 13109 defined nucleotide sequences given in the
 (or complements/ fragments). The probe hybridises at high
 a nucleic acid molecule expressed in the human adult liver.
 d for predicting, measuring and displaying gene expression
 ived from human adult liver. The genes identified may be
 netic liver diseases such as cirrhosis,
 naemia, hyperlipidaemia and hypercholesterolaemia which is
 h coronary heart disease. ABG47348-ABG59930 represent human
 on encoded peptides of the invention. Note: The sequence
 r this patent does not appear in the printed specification
 ed in electronic format directly from WIPO at
 ub/published_pct_sequences

2.8%; Score 8; DB 4; Length 65;
 100.0%; Pred. No. 27;
 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 LGLGLALA 68
 |||||
 Db 22 LGLGLALA 29

RESULT 41
 ABG46737
 ID ABG46737 standard; peptide; 65 AA.
 XX
 AC ABG46737;
 XX
 DT 19-AUG-2002 (first entry)
 XX
 DE
 XX
 KW Human peptide encoded by genome-derived single exon probe SEQ ID
 KW Human; single exon probe; asthma; lung cancer; COPD; ILD;
 KW chronic obstructive pulmonary disease; interstitial lung disease;
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
 KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndr
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
 KW Primary ciliary dyskinesia; pulmonary hypertension;
 KW hyaline membrane disease.
 XX
 OS Homo sapiens.
 XX
 PN WO200186003-A2.
 XX
 PD 15-NOV-2001.
 XX
 PF 30-JAN-2001; 2001WO-US000665.
 XX
 PR 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00608408.
 PR 03-AUG-2000; 2000US-00632366.
 PR 21-SEP-2000; 2000US-0234587P.
 PR 27-SEP-2000; 2000US-0236359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 DR WPI; 2002-114183/15.
 XX
 PT Spatially-addressable set of single exon nucleic acid probes, used
 PT measure gene expression in human lung samples.
 XX
 PS Claim 27; SEQ ID NO 36402; 634pp; English.
 XX
 CC The invention relates to a spatially-addressable set of single exc
 CC nucleic acid probes for measuring gene expression in a sample deri
 CC from human lung comprising single exon nucleic acid probes having
 CC 12614 nucleic acid sequences mentioned in the specification, or th
 CC complements or the 12387 open reading frames derived from the 1261
 CC probes. Also included are a microarray comprising the novel set of
 CC : the novel set of probes which hybridise at high stringency to a
 CC acid expressed in the human lung; measuring gene expression in a s
 CC derived from human lung, comprising (a) contacting the array with
 CC collection of detectably labeled nucleic acids derived from human
 CC mRNA, and (b) measuring the label detectably bound to each probe o
 CC array; identifying exons in a eukaryotic genome, comprising (a)
 CC algorithmically predicting at least one exon from genomic sequence
 CC the eukaryote; and (b) detecting specific hybridisation of detecta
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon p
 CC having a fragment identical to the predicted exon, the probe is in
 CC in the above mentioned microarray; assigning exons to a single gen
 CC comprising (a) identifying exons from genomic sequence by the meth
 CC above and (b) measuring the expression of each of the exons in sev
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern o

the exons in the tissues and/or cell types indicates that could be assigned to a single gene; a peptide comprising one or more amino acids, mentioned in the specification, or encoded by the reading frames (ORF). The probes are used for gene expression analysis for identifying exons in a gene, particularly using human cDNA and for the study of lung diseases such as asthma, lung cancer, obstructive pulmonary disease (COPD), interstitial lung disease, familial idiopathic pulmonary fibrosis, neurofibromatosis, sarcoidosis, Gaucher's disease, Niemann-Pick disease, Hermansky-Raudon, lymphangioleiomyomatosis, pulmonary alveolar proteinosis, idiopathic pulmonary hypertension and hyaline membrane disease. The probe is a peptide/protein encoded by a single exon probe of the gene. Note: The sequence data for this patent did not form part of the specification, but was obtained in electronic format from WIPO at ftp.wipo.int/pub/published_pct_sequences

Query Match 2.8%; Score 8; DB 5; Length 65;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 0; Mismatches 0; Indels 0; Gaps 0;
Conservative 0; Mismatches 0; Indels 0; Gaps 0;
HLALA 68
|||||
HLALA 29

Standard; protein; 69 AA.

(first entry)

Diagnostic protein #3714.

Some mapping; gene mapping; gene therapy; forensic;
nt; medical imaging; diagnostic; genetic disorder.

2.

2001WO-US008631.

2000US-00540217.

2000US-00649167.

INC.

Liu C, Tang YT;

362/73.

910.

polynucleotide and encoded polypeptides, useful in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits and to assess

ID NO 34082; 103pp; English.

relates to isolated polynucleotide (I) and polypeptide (II)
is useful as hybridisation probes, polymerase chain
primers, oligomers, and for chromosome and gene mapping,
inant production of (II). The polynucleotides are also used
as expressed sequence tags for identifying expressed
useful in gene therapy techniques to restore normal

activity of (II) or to treat disease states involving (II). (II)
useful for generating antibodies against it, detecting or quantifying
polypeptide in tissue, as molecular weight markers and as a food
supplement. (II) and its binding partners are useful in medical
of sites expressing (II). (I) and (II) are useful for treating d
involving aberrant protein expression or biological activity. Th
polypeptide and polynucleotide sequences have applications in
diagnostics, forensics, gene mapping, identification of mutation
responsible for genetic disorders or other traits to assess biod
and to produce other types of data and products dependent on DNA
amino acid sequences. ABG00010-ABG30377 represent novel human di
amino acid sequences of the invention. Note: The sequence data f
patent did not appear in the printed specification, but was obta
electronic format directly from WIPO at
ftp.wipo.int/pub/published_pct_sequences

Sequence 69 AA;

Query Match 2.8%; Score 8; DB 4; Length 69;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

Oy 61 LGLGLALA 68
Db 16 LGLGLALA 23
|||||

RESULT 43

ABG03663

ID ABG03663 standard; protein; 71 AA.

AC ABG03663;

DT 13-FEB-2002 (first entry)

DE Novel human diagnostic protein #3654.

Human; chromosome mapping; gene mapping; gene therapy; forensic;
food supplement; medical imaging; diagnostic; genetic disorder.

OS Homo sapiens.

XX WO200175067-A2.

PN 11-OCT-2001.

PD 30-MAR-2001; 2001WO-US008631.

PF 31-MAR-2000; 2000US-00540217.

PR 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

DR N-PSDB; AAS67850.

XX New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity.

PS Claim 20; SEQ ID NO 34022; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypept
sequences. (I) is useful as hybridisation probes, polymerase chai
reaction (PCR) primers, oligomers, and for chromosome and gene ma
in and recombinant production of (II). The polynucleotides are al
in diagnostics as expressed sequence tags for identifying express
genes. (I) is useful in gene therapy techniques to restore normal
activity of (II) or to treat disease states involving (II). (II)
useful for generating antibodies against it, detecting or quantit

n tissue, as molecular weight markers and as a food
 (II) and its binding partners are useful in medical imaging
 (II). (I) and (II) are useful for treating disorders
 of protein expression or biological activity. The
 and polynucleotide sequences have applications in
 forensics, gene mapping, identification of mutations
 or genetic disorders or other traits to assess biodiversity
 a other types of data and products dependent on DNA and
 sequences. ABG00010-ABG30377 represent novel human diagnostic
 sequences of the invention. Note: The sequence data for this
 appear in the printed specification, but was obtained in
 mat directly from WIPO at
 pub/published_pct_sequences

A;
 2.8%; Score 8; DB 4; Length 71;
 arity 100.0%; Pred. No. 29; 0; Indels 0; Gaps 0;
 nservative 0; Mismatches 0; Mismatches 0; Indels 0; Gaps 0;
 ALA 69
 |||||
 ALA 23

lard; protein; 84 AA.

(first entry)

ium acnes immunogenic protein #22178.

; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
 hthralmitis; bone; joint; central nervous system; ELISA;
 esion; acne vulgaris; enzyme linked immunosorbent assay;
 ; osteopathic; neuroprotectant.

ium acnes.

001WO-US012865.

000US-0199047P.

000US-0208841P.

000US-0216747P.

CORP.

ersing DH, Mitcham JL, Wang SS, Bhatia A;
 J, Zhang Y, Jen S, Carter D;

74/71.

15.

ium acnes polypeptides and nucleic acids useful for
 ainst and diagnosing infections, especially useful for
 vulgaris.

ID NO 22477; 1069pp; English.

910S-AAU68017 represent Propionibacterium acnes immunogenic
 The proteins and their associated DNA sequences are used in
 prevention and diagnosis of medical conditions caused by
 disorders include SAPHO syndrome (synovitis, acne,
 perosis and osteomyelitis), uveitis and endophthalmitis.
 so involved in infections of bone, joints and the central
 , however it is particularly involved in the inflammatory

lesions associated with acne vulgaris. A method for detecting the
 presence or absence of P. acnes in a patient comprises contacting
 sample with a binding agent that binds to the proteins of the in
 and determining the amount of bound protein in the sample. The
 polypeptides may be used as antigens in the production of antibod
 specific for P. acnes proteins. These antibodies can be used to
 downregulate expression and activity of P. acnes polypeptides and
 therefore treat P. acnes infections. The antibodies may also be u
 diagnostic agents for determining P. acnes presence, for example,
 enzyme linked immunosorbent assay (ELISA). Note: The sequence dat
 this patent did not form part of the printed specification, but w
 obtained in electronic format directly from WIPO at
 ftp.wipo.int/pub/published_pct_sequences

SQ Sequence 84 AA;

Query Match 2.8%; Score 8; DB 4; Length 84;
 Best Local Similarity 100.0%; Pred. No. 34;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 259 LRITLPM 266

Db 34 LRITLPM 41

RESULT 45

ABM57801

ID ABM57801 standard; protein; 84 AA.

AC ABM57801;

DT 20-OCT-2003 (first entry)

DE Propionibacterium acnes predicted ORF-encoded polypeptide #22477.

KW Acne vulgaris; antiseborrheic; dermatological; antibacterial;
 immunostimulant; immune response; vaccine.

OS Propionibacterium acnes.

PN WO2003033515-A1.

PD 24-APR-2003.

PF 11-OCT-2002; 2002WO-US032727.

PR 15-OCT-2001; 2001US-00978825.

XX (CORI-) CORIXA CORP.

PI Mitcham JL, Skeiky YAW, Persing DH, Bhatia A, Maisonneuve JL;
 PI Zhang Y, Wang S, Jen S, Lodes MJ, Benson DR, Jones R, Carter
 PI Barth B, Vallieve-Douglas J;

XX WPI; 2003-381789/36.

DR N-PSDB; ACF64544.

XX New Propionibacterium acnes polypeptides and polynucleotides encod
 PT polypeptide, useful for diagnosing, preventing or treating acne v
 PT or for stimulating an immune response specific for a P. acnes prot
 PS Example 1; SEQ ID NO 22477; 1481pp; English.

XX The invention relates to an isolated polynucleotide (ACF64435-ACF6
 CC encoding a Propionibacterium acnes protein. The invention also rel
 CC polypeptides encoded by the polynucleotides (ABM35624-ABM64536) an
 CC immunogenic fragments of P. acnes polypeptides. The invention
 CC additionally encompasses expression vectors and host cells compris
 CC polynucleotide of the invention; antibodies against polypeptides c
 CC invention; fusion proteins comprising a polypeptide of the inventi
 CC method for stimulating an immune response specific for a P. acnes
 CC polypeptide and an isolated T cell population comprising T cells p
 CC via this method; a vaccine composition (comprising P. acnes polype

ies, antibodies, fusion proteins, T cell populations, or
 enting cells that express the polypeptide); a method and kit
 g or determining the presence or absence of P. acnes in a
 method for inhibiting the development of P. acnes in a
 P. acnes polypeptides, polynucleotides, antibodies, fusion
 cell populations or antigen-presenting cells that express the
 are useful for diagnosing, preventing or treating acne
 for stimulating an immune response specific for a P. acnes
 polynucleotides can also be used as probes or primers for
 hybridisation. The vaccine composition is useful for the
 of an immune response against P. acnes, or for treating acne,
 is useful for performing a diagnostic assay. The present
 resents a polypeptide predicted to be encoded by an ORF (open
 e) contained within the P. acnes polynucleotides of the
 te: The sequence data for this patent did not form part of
 specification, but was obtained in electronic format directly
 ftp.wipo.int/pub/published_pct_sequences

AA;

2.8%; Score 8; DB 6; Length 84;
 larity 100.0%; Pred. No. 34;
 Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RTLPW 266

|||||
 RTLPW 41

ard; protein; 110 AA.

(first entry)

diagnostic protein #20250.

some mapping; gene mapping; gene therapy; forensic;
 nt; medical imaging; diagnostic; genetic disorder.

12.

2001WO-US008631.

2000US-00540217.

2000US-00649167.

INC.

Liu C, Tang YT;

1362/73.

446.

polynucleotide and encoded polypeptides, useful in
 forensics, gene mapping, identification of mutations
 or genetic disorders or other traits and to assess

ID NO 50618; 103pp; English.

relates to isolated polynucleotide (I) and polypeptide (II)
 is useful as hybridisation probes, polymerase chain
 primers, oligomers, and for chromosome and gene mapping,
 nant production of (II). The polynucleotides are also used
 as expressed sequence tags for identifying expressed
 useful in gene therapy techniques to restore normal

CC activity of (II) or to treat disease states involving (II). (II)
 CC useful for generating antibodies against it, detecting or quanti
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical
 CC of sites expressing (II). (I) and (II) are useful for treating d
 CC involving aberrant protein expression or biological activity. Th
 CC polypeptide and polynucleotide sequences have application in
 CC diagnostics, forensics, gene mapping, identification of mutation
 CC responsible for genetic disorders or other traits to assess blood
 CC and to produce other types of data and products dependent on DNA
 CC amino acid sequences. ABG00010-ABG30377 represent novel human di
 CC amino acid sequences of the invention. Note: The sequence data f
 CC patent did not appear in the printed specification, but was obta
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 110 AA;

Query Match 2.8%; Score 8; DB 4; Length 110;
 Best Local Similarity 100.0%; Pred. No. 45;
 Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 61 LGLGLALA 68

Db 44 LGLGLALA 51

RESULT 47

AAO08094

ID AAO08094 standard; protein; 117 AA.

XX AAO08094;

XX AAO08094;

DT 06-NOV-2001 (first entry)

DE Human polypeptide SEQ ID NO 21986.

XX

Human; cytokine; cell proliferation; cell differentiation; gene
 vaccine; peptide therapy; stem cell growth factor; haematopoiesi
 tissue growth factor; immunomodulatory; cancer; leukaemia;
 nervous system disorders; arthritis; inflammation.

OS Homo sapiens.

XX WO200164835-A2.

XX 07-SEP-2001.

XX 26-FEB-2001; 2001WO-US004927.

XX 28-FEB-2000; 2000US-00515126.

XX 18-MAY-2000; 2000US-00577409.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-514838/56.

XX N-PSDB; AAI88025.

XX Isolated nucleic acids and polypeptides, useful for preventing d:

XX and treating e.g. leukemia, inflammation and immune disorders.

XX Claim 20; SEQ ID NO 21986; 1399pp + Sequence Listing; English.

XX The invention relates to human polynucleotides (AAI79941-AAI93841)

XX the encoded proteins (AAO00010-AAO13910) that exhibit activity el

XX cytokine, cell proliferation or cell differentiation or which may

XX production of other cytokines in other cell populations. The

XX polynucleotides and polypeptides are useful in gene therapy, vac

XX peptide therapy. The polypeptides have various cytokine-like acti

XX e.g. stem cell growth factor activity, haematopoiesis regulating

XX activity, tissue growth factor activity, immunomodulatory activit

in activity and may be useful in the diagnosis and/or cancer, leukaemia, nervous system disorders, arthritis and Note: The sequence data for this patent did not form part of the specification, but was obtained in electronic format from WIPO at ftp.wipo.int/pub/published_pct_sequences

AA;

2.8%; Score 8; DB 4; Length 117;
arity 100.0%; Pred. No. 47;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

ALA 68
||||
ALA 61

ard; protein; 184 AA.

(first entry)

lanogaster polypeptide SEQ ID NO 29928.

velopmental biology; cell signalling; insecticide;

lanogaster.

2.

2001WO-US009231.

2000US-0191637P.

2000US-00614150.

ip NY.

lams M, Li PWD, Myers EW;

60/75.

115.

nucleic acid detection reagent for detecting 1000 or more
ophila and for elucidating cell signalling and cell-cell

Q ID NO 29928; 21pp + Sequence Listing; English.

relates to an isolated nucleic acid detection reagent
acting 1000 or more genes from Drosophila. The invention is
developmental biology and in elucidating cell signalling and
fractions in higher eukaryotes for the development of
therapeutics and pharmaceutical drugs. The invention
mic DNA sequences (ABL16176-ABL30511), expressed DNA
01840-ABL16175) and the encoded proteins (AB57737-
sequence data for this patent did not form part of the
ication, but was obtained in electronic format directly
ftp.wipo.int/pub/published_pct_sequences

AA;

2.8%; Score 8; DB 4; Length 184;
arity 100.0%; Pred. No. 73;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

WSL 79
|||

Db 16 LLLAVVSL 23

RESULT 49

ID ABP28041 standard; protein; 190 AA.

XX ABP28041;

XX AC ABP28041;

XX DT 02-JUL-2002 (first entry)

XX DE Streptococcus polypeptide SEQ ID NO 5258.

XX KW Streptococcus; GAS; GBS; group B streptococcus; Streptococcus aga

XX KW Group A streptococcus; Streptococcus pyogenes; antibacterial;

XX KW antiinflammatory; infection; vaccine; meningitis; gene therapy.

XX OS Streptococcus agalactiae.

XX PN WO200234771-A2.

XX PD 02-MAY-2002.

XX PF 29-OCT-2001; 2001WO-GB004789.

XX PR 27-OCT-2000; 2000GB-00026333.

XX PR 24-NOV-2000; 2000GB-00028727.

XX PR 07-MAR-2001; 2001GB-00005640.

XX PA (CHIR-) CHIRON SPA.

XX PA (GENO-) INST GENOMIC RES.

XX PI Telford J, Masighani V, Margarit Y RosI, Grandi G, Fraser C;

XX PI Tettelin H;

XX DR WPI; 2002-352536/38.

XX DR N-PSDB; ABN68672.

XX PT New Streptococcus protein for the treatment or prevention of infe

XX PT disease caused by Streptococcus bacteria, such as meningitis, and

XX PT detecting a compound that binds to the protein.

XX PS Claim 1; Page 3689; 4525pp; English.

XX CC The invention relates to a protein (ABP25413-ABP30895) from group

XX CC Streptococcus/GBS (Streptococcus agalactiae) or group A streptoco

XX CC (Streptococcus pyogenes), comprising one of 5483 sequences (S1),

XX CC the specification. The proteins have antibacterial and antinflam

XX CC activity. (I), nucleic acids encoding (I), ABN66044-ABN71526 and

XX CC antibodies that bind (I) are used in the manufacture of medicamen

XX CC the treatment or prevention of infection or disease caused by

XX CC Streptococcus bacteria, particularly S. agalactiae and S. pyogen

XX CC Nucleic acids encoding (I) are used to detect Streptococcus in a

XX CC biological sample. (I) is used to determine whether a compound bin

XX CC (I). A composition comprising (I) or a nucleic acid encoding (I),

XX CC used as a vaccine or diagnostic composition. The disease caused by

XX CC Streptococcus that is prevented or treated may be meningitis. Nuc

XX CC acid encoding (I) may be used to recombinantly produce (I) and ma

XX CC used in gene therapy. Antibodies to (I) are used for affinity

XX CC chromatography, immunoassays, and distinguishing/identifying

XX CC Streptococcus proteins

XX SQ Sequence 190 AA;

Query Match 2.8%; Score 8; DB 5; Length 190;
Best Local Similarity 100.0%; Pred. No. 75;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps

QY 258 SLRIRLTP 265

Db 50 SLRIRLTP 57

ward; protein; 198 AA.

(first entry)

oded protein SEQ ID NO: 1209.

y; dog; fruit fly; Yeast; hamster; macaque; horse;
forensic test; Gene mapping; genetic disorder; Biodiversity;
nutrition.

2.

2001WO-US002687.

2000US-00491404.

2000US-00617746.

2000US-00631451.

2000US-00663870.

INC.

C, Zhou P, Qian XB, Wang Z, Chen R, Asundi V;
ac RA, Zhang J, Werhman T;
164/51.
343.

Peptide for treatment of diseases, diagnostics, raising
id research use.

873; 1275pp; English.

vention provides the protein and coding sequences of novel
a variety of organisms, including human, dog, cat, horse,
ater, monkey, macaque, yeast, bacteria, fruit fly, sea
mato. These were derived from expressed sequence tags (ESTs)
nism of interest. They can be used in diagnostics,
ne mapping, identification of mutations, to assess
and for nutritional purposes. The present sequence is a
e invention

AA;

2.8%; Score 8; DB 4; Length 198;
arity 100.0%; Pred. No. 78;
onservative 0; Mismatches 0; Indels 0; Gaps 0;

PLAL 61
||||
PLAL 9

ward; protein; 222 AA.

(first entry)

antigen HPAMG11, SEQ ID NO:2806.

an antigen; ovary; ovarian; breast; cancer; tumour;
r; breast cancer; tumour; reproductive system disorder;

infertility; pregnancy disorder; anovulation; polycystic ovary s
PCOS; ovarian cyst; dysmenorrhoea; endocrine disorder; infection
inflammatory condition; immune disorder; blood disorder;
cardiovascular disorder; respiratory disorder; neurological diso
gastrointestinal disorder; urinary system disorder; drug screeni
gene therapy; chromosome mapping; forensic analysis;
antibody preparation; cytostatic; immunomodulatory; neuroprotect
antiinflammatory; gynaecological; reproductive.

Homo sapiens.

WO200200677-A1.

03-JAN-2002.

07-JUN-2001; 2001WO-US018569.

07-JUN-2000; 2000US-0209467P.

(HUMA-) HUMAN GENOME SCI INC.

Birse CE, Rosen CA;

WPI; 2002-147878/19.

N-PSDB; ABQ54751.

Isolated nucleic acid molecules encoding novel ovarian polypeptid
useful in the prevention, treatment and diagnosis of cancer (e.g
cancer), immune disorders, cardiovascular disorders and neurolog
diseases.

Claim 11; SEQ ID NO 2806; 2922pp; English.

The invention relates to 2175 novel human ovarian antigens (ABP4
ABP43228) and to cDNAs encoding them (ABQ54131-ABQ56305), and al
encapsulates polypeptides 90% identical and polynucleotides 95% i
to the sequences of the invention. The invention additionally re
recombinant vectors and host cells comprising human ovarian anti
polynucleotides, antibodies against human ovarian antigens, and
of ovarian antigen polynucleotides and polypeptides in diagnosis
treating, prophosing or preventing various ovary and/or breast-
disorders. Such conditions include ovarian cancer and breast can
metastatic tumours of ovarian or breast origin, reproductive sys
disorders (e.g., infertility, disorders of pregnancy, anovulation
polycystic ovary syndrome, ovarian cysts, and dysmenorrhoea), en
disorders, infections (e.g., chlamydia, HIV, toxoplasmosis, and
shock syndrome), inflammatory conditions (e.g., mastitis, ophor
vaginits), immune disorders (e.g., congenital and acquired
immunodeficiencies, autoimmune ophoritis, systemic lupus erythe
blood-related disorders (e.g., anaemia), cardiovascular disorder
respiratory disorders, neurological disorders, gastrointestinal
and urinary system disorders. Ovarian antigen polypeptides and
polynucleotides may also be used in screening for compounds whic
modulate ovarian antigen expression or activity. The polynucleot
further be used for gene therapy, chromosome mapping, in the
identification of individuals and in forensic analysis, and the
polypeptides may be used as food additives or to prepare antibodi
useful in disease diagnosis, drug targeting and phenotyping. The
sequence represents a human ovarian antigen of the invention. Not
sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly fr
at ftp.wipo.int/pub/published_pct_sequences

Sequence 222 AA;

Query Match 2.8%; Score 8; DB 5; Length 222;
Best Local Similarity 100.0%; Pred. No. 87;
Matches 8; Conservative 0; Mismatches 0; Indels 0; C

QY 60 ALGLGLAL 67
|||||

DB 190 ALGLGLAL 197

dard; protein; 286 AA.
 (first entry)
 agnostic protein #16270.
 some mapping; gene mapping; gene therapy; forensic;
 nt; medical imaging; diagnostic; genetic disorder.
 2.
 2001WO-US008631.
 2000US-00540217.
 2000US-00649167.
 INC.
 Liu C, Tang YT;
 362/73.
 166.
 polynucleotide and encoded polypeptides, useful in
 forensics, gene mapping, identification of mutations
 or genetic disorders or other traits and to assess
 ID NO 46638; 103pp; English.
 relates to isolated polynucleotide (I) and polypeptide (II)
 is useful as hybridisation probes, polymerase chain
 reaction (PCR) primers, oligomers, and for chromosome and gene ma
 inant production of (II). The polynucleotides are also used
 as expressed sequence tags for identifying expressed
 useful in gene therapy techniques to restore normal
 (II) or to treat disease states involving (II). (II) is
 useful for generating antibodies against it, detecting or quantitat
 ing tissue, as molecular weight markers and as a food
 supplement. (II) and its binding partners are useful in medical i
 maging (II). (I) and (II) are useful for treating disorders
 involving aberrant protein expression or biological activity. The
 id polynucleotide sequences have applications in
 forensics, gene mapping, identification of mutations
 or genetic disorders or other traits to assess biodiversity
 other types of data and products dependent on DNA and
 sequences. ABG0010-ABG30377 represent novel human dia
 gnoses of the invention. Note: The sequence data for this
 appear in the printed specification, but was obtained in
 mat directly from WIPO at
 ftp.wipo.int/pub/published_pct_sequences
 Query Match 2.8%; Score 8; DB 4; Length 286;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 0; Mismatches 0; Indels 0; Gaps 0;
 Conservative 0;
 VSLGSRAS 258
 VSLGSRAS 76
 ABG01186
 ID ABG01186 standard; protein; 307 AA.
 AC ABG01186;
 DT 13-FEB-2002 (first entry)
 DE Novel human diagnostic protein #1177.
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder.
 OS Homo sapiens.
 PN WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US008631.
 XX 31-MAR-2000; 2000US-00540217.
 PR 23-AUG-2000; 2000US-00649167.
 XX (HYSE-) HYSEQ INC.
 PA Drmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 DR N-PSDB; AAS65373.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX Claim 20; SEQ ID NO 31545; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and polypept
 sequences. (I) is useful as hybridisation probes, polymerase chain
 reaction (PCR) primers, oligomers, and for chromosome and gene ma
 and in recombinant production of (II). The polynucleotides are al
 in diagnostics as expressed sequence tags for identifying expres
 genes. (I) is useful in gene therapy techniques to restore normal
 activity of (II) or to treat disease states involving (II). (II)
 useful for generating antibodies against it, detecting or quantit
 polypeptide in tissue, as molecular weight markers and as a food
 supplement. (II) and its binding partners are useful in medical i
 of sites expressing (II). (I) and (II) are useful for treating di
 involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodi
 CC and to produce other types of data and products dependent on DNA
 CC amino acid sequences. ABG0010-ABG30377 represent novel human dia
 CC amino acid sequences of the invention. Note: The sequence data fo
 CC patent did not appear in the printed specification, but was obtain
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX Sequence 307 AA;
 SQ
 Query Match 2.8%; Score 8; DB 4; Length 307;
 Best Local Similarity 100.0%; Pred. No. 1.2e+02;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G
 Conservative 0;
 VSLGSRAS 84
 VSLGSRAS 10
 RESULT 54
 ADB79952
 ID ADB79952 standard; protein; 342 AA.

QY
DE
EE
AE
IE
XX
AC
XX
AT
DT
XX
DE
XX
KW
KW
XX
XX
OS
XX
PN
XX
PG
XX
PF
XX
PR
XX
PR

(first entry)

BB 1 progression enhanced protein, SEQ ID 192.

ain; streptozocin-induced diabetes; rat.

gicus.

2002EP-00255249.

2001GB-00018354.

2002GB-00002910.

ER LAMBERT CO.

A, Dixon AK, Lee K, Pinnock RD;

5407/38.

9953.

ed gene sequences and encoded polypeptides that are
in the spinal cord in response to streptozocin-induced
screening compounds for the treatment of pain, or for
ain.

326-327; 334pp; English.

vention relates to nucleotide sequences which are useful in
of compounds for the treatment of pain, or for the
pain. The nucleotide sequences are up-regulated in the
in response to streptozocin-induced diabetes. The present
used to illustrate the invention.

AA;

2.8%; Score 8; DB 7; Length 342;

arity 100.0%; Pred. No. 1.3e+02;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GLAL 67

||||

GLAL 317

dard; protein; 370 AA.

(first entry)

diagnostic protein #13382.

ome mapping; gene mapping; gene therapy; forensic;
nt; medical imaging; diagnostic; genetic disorder.

12.

2001WO-US008631.

2000US-00540217.

PR 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSRQ INC.

XX Drmanac RT, Liu C, Tang YT;

PI WPI; 2001-639362/73.

XX DR N-PSDB; AAS7578.

XX

New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutation
responsible for genetic disorders or other traits and to assess
biodiversity.

Claim 20; SEQ ID NO 43750; 103pp; English.

The invention relates to isolated polynucleotide (I) and polypep
sequences. (I) is useful as hybridisation probes, polymerase cha
reaction (PCR) primers, oligomers, and for chromosome and gene m
and in recombinant production of (II). The polynucleotides are a
in diagnostics as expressed sequence tags for identifying expres
genes. (I) is useful in gene therapy techniques to restore norma
activity of (II) or to treat disease states involving (II). (II)
useful for generating antibodies against it, detecting or quanti
polypeptide in tissue, as molecular weight markers and as a food
supplement. (II) and its binding partners are useful in medical
of sites expressing (II). (I) and (II) are useful for treating d
involving aberrant protein expression or biological activity. Th
polypeptide and polynucleotide sequences have application in
diagnostics, forensics, gene mapping, identification of mutation
responsible for genetic disorders or other traits to assess biod
and to produce other types of data and products dependent on DNA
amino acid sequences. ABG00010-ABG30377 represent novel human di
patent did not appear in the invention. Note: The sequence data f
electronic format directly from WIPO at
ftp.wipo.int/pub/published_pct_sequences

XX Sequence 370 AA;

Query Match

Best Local Similarity 2.8%; Score 8; DB 4; Length 370;

Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 63 LGALACL 70

||||||

Db 28 LGALACL 35

RESULT 56

ABG05012

ID ABG05012 standard; protein; 370 AA.

XX AC

XX ABG05012;

XX DT

XX 13-FEB-2002 (first entry)

XX DE

XX Novel human diagnostic protein #5003.

XX KW

Human; chromosome mapping; gene mapping; gene therapy; forensic;
food supplement; medical imaging; diagnostic; genetic disorder.

XX OS

XX Homo sapiens.

XX PN

XX WO200175067-A2.

XX PD

XX 11-OCT-2001.

XX PF

XX 30-MAR-2001; 2001WO-US008631.

XX PR

XX 31-MAR-2000; 2000US-00540217.

XX PR

XX 23-AUG-2000; 2000US-00649167.

XX XX

INC.

Liu C, Tang YT;

362/73.
199.

polynucleotide and encoded polypeptides, useful in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits and to assess

ID NO 35371; 103pp; English.

relates to isolated polynucleotide (I) and polypeptide (II)
is useful as hybridisation probes, polymerase chain
primers, oligomers, and for chromosome and gene map-
ing and in recombinant production of (II). The polynucleotides are also used
as expressed sequence tags for identifying expressed
genes. (I) is useful in gene therapy techniques to restore normal
activity of (II) or to treat disease states involving (II). (II) is
useful for generating antibodies against it, detecting or quantitat-
ing polypeptide in tissue, as molecular weight markers and as a food
supplement. (II) and its binding partners are useful in medical
diagnostics, forensics, gene mapping, identification of mutations
in polynucleotide sequences have applications in
other types of data and products dependent on DNA and
amino acid sequences. ABG0010-ABG30377 represent novel human dia-
gnostic sequences of the invention. Note: The sequence data fo-
rmat directly from WIPO at
ftp://published_pct_sequences

A;

Query Match 2.8%; Score 8; DB 4; Length 370;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 0; Mismatches 0; Indels 0; Gaps 0;

ACL 70
|||
ACL 35

ard; protein; 370 AA.

first entry)

agnostic protein #18106.

ome mapping; gene mapping; gene therapy; forensic;
t; medical imaging; diagnostic; genetic disorder.

001WO-US008631.

000US-00540217.

000US-00649167.

INC.

PI Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.
DR N-PSDB; AAS82302.

XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
XX biodiversity.

Claim 20; SEQ ID NO 48474; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypept
CC sequences (II) is useful as hybridisation probes, polymerase chai
CC reaction (PCR) primers, oligomers, and for chromosome and gene ma
CC and in recombinant production of (II). The polynucleotides are al
CC in diagnostics as expressed sequence tags for identifying express
CC genes. (I) is useful in gene therapy techniques to restore normal
CC activity of (II) or to treat disease states involving (II). (II)
CC useful for generating antibodies against it, detecting or quantitat
CC polypeptide in tissue, as molecular weight markers and as a food
CC supplement. (II) and its binding partners are useful in medical
CC of sites expressing (II). (I) and (II) are useful for treating di
CC involving aberrant protein expression or biological activity. The
CC polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodi
CC and to produce other types of data and products dependent on DNA
CC amino acid sequences. ABG0010-ABG30377 represent novel human dia
CC amino acid sequences of the invention. Note: The sequence data fo
CC patent did not appear in the printed specification, but was obtain
CC electronic format directly from WIPO at
CC ftp://published_pct_sequences

XX Sequence 370 AA;

Query Match 2.8%; Score 8; DB 4; Length 370;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 63 LGLALACL 70
|||
Db 28 LGLALACL 35

RESULT 58
ABU36537

ID ABU36537 standard; protein; 372 AA.

XX AC ABU36537;

XX DT 19-JUN-2003 (first entry)

DE Protein encoded by Prokaryotic essential gene #22064.

XX Antisense; prokaryotic essential gene; cell proliferation; drug de
XX Mycobacterium tuberculosis.

XX PN WO200277183-A2.

XX PD 03-OCT-2002.

XX PF 21-MAR-2002; 2002WO-US009107.

XX PR 21-MAR-2001; 2001US-00815242.

XX PR 06-SEP-2001; 2001US-00948993.

XX PR 25-OCT-2001; 2001US-0342923P.

XX PR 08-FEB-2002; 2002US-00072851.

XX PR 06-MAR-2002; 2002US-0362699P.

XX PA (ELIT-) ELITRA PHARM INC.
XX

radio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW,
 vick JD, Carr GU, Yamamoto R, Forsyth RA, Xu HH;
 3926/02.
 3926/02.
 a nucleic acids, useful for identifying proteins or screening
 as nucleic acids required for cellular proliferation to
 idate molecules for rational drug discovery programs.
) ID NO 64461; 1766pp; English.
 i relates to an isolated nucleic acid comprising any one of
 sense sequences given in the specification where expression
 c acid inhibits proliferation of a cell. Also included are:
 comprising a promoter operably linked to the nucleic acid
 peptide whose expression is inhibited by the antisense
 (2) a host cell containing the vector; (3) an isolated
 or its fragment whose expression is inhibited by the
 leic acid; (4) an antibody capable of specifically binding
 de; (5) producing the polypeptide; (6) inhibiting cellular
 i or the activity of a gene in an operon required for
 i; (7) identifying a compound that influences the activity of
 duct or that has an activity against a biological pathway
 proliferation, or that inhibits cellular proliferation; (8)
 a gene required for cellular proliferation or the biological
 which a proliferation-required gene or its gene product lies
 which the test compound that inhibits proliferation of an
 ; (9) manufacturing an antibiotic; (10) profiling a
 ; (11) a culture comprising strains in which the gene
 repressed or underexpressed; (12) determining the extent
 of the strains is present in a culture or collection of
 13) identifying the target of a compound that inhibits the
 of an organism. The antisense nucleic acids are useful for
 proteins or screening for homologous nucleic acids required
 proliferation to isolate candidate molecules for rational
 y programs, or for screening homologous nucleic acids
 proliferation in cells other than *S. aureus*, *S. typhimurium*,
 or *P. aeruginosa*. The present sequence is encoded by one of
 okaryotic essential genes. Note: The sequence data for this
 t form part of the printed specification, but was obtained
 format directly from WIPO at
 pub/published_pct_sequences

AA;

2.8%; Score 8; DB 6; Length 372;
 arity 100.0%; Pred. No. 1.4e+02;
 onservative 0; Mismatches 0; Indels 0; Gaps 0;

LALA 68
 ||||
 LALA 103

dard; protein; 424 AA.

(first entry)

diagnostic protein #15604.

some mapping; gene mapping; gene therapy; forensic;
 nt; medical imaging; diagnostic; genetic disorder.

2.

XX 30-MAR-2001; 2001WO-US008631.
 XX 31-MAR-2000; 2000US-00540217.
 PR 23-AUG-2000; 2000US-00649167.
 XX (HYSE-) HYSEQ INC.
 PA Drmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 DR N-PSDB; AAS79800.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutation;
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX Claim 20; SEQ ID NO 45972; 103pp; English.
 PS The invention relates to isolated polynucleotide (I) and polypep
 CC sequences. (I) is useful as hybridisation probes, polymerase cha
 CC reaction (PCR) primers, oligomers, and for chromosome and gene m
 CC and in recombinant production of (II). The polynucleotides are a
 CC in diagnostics as expressed sequence tags for identifying expres
 CC genes. (II) is useful in gene therapy techniques to restore norma
 CC activity of (II) or to treat disease states involving (II). (II)
 CC useful for generating antibodies against it, detecting or quantifi
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical
 CC of sites expressing (II). (I) and (II) are useful for treating d
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutation
 CC responsible for genetic disorders or other traits to assess biod
 CC and to produce other types of data and products dependent on DNA
 CC amino acid sequences. ABG00010-ABG30377 represent novel human di
 CC amino acid sequences of the invention. Note: The sequence data f
 CC patent did not appear in the printed specification, but was obta
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX Sequence 424 AA;

Query Match 2.8%; Score 8; DB 4; Length 424;
 Best Local Similarity 100.0%; Pred. No. 1.6e+02;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; G

Qy 60 ALGGLAL 67

Db 392 ALGGLAL 399

RESULT 60

AAM23752

ID AAM23752 standard; protein; 430 AA.

XX AAM23752;

DT 12-OCT-2001 (first entry)

DE Human EST encoded protein SEQ ID NO: 1277.

XX Human; sheep; pig; cow; fruit fly; yeast; hamster; macaque; horse
 KW tomato; monkey; dog; sea urchin; expressed sequence tag; EST;
 KW diagnostics; forensic test; gene mapping; genetic disorder; biodi
 KW gene therapy; nutrition.

OS Homo sapiens.

XX WO200154477-A2.

XX 02-AUG-2001.

2001WO-US002687.

2000US-00491404.

2000US-00617746.

2000US-00631451.

2000US-00663870.

INC.

C, Zhou P, Qian XB, Wang Z, Chen R, Asundi V;
ac RA, Zhang J, Werhman T;

164/51.

411.

peptide for treatment of diseases, diagnostics, raising
a research use.

920; 1275pp; English.

vention provides the protein and coding sequences of novel
a variety of organisms, including human, dog, cat, horse,
ster, monkey, macaque, yeast, bacteria, fruit fly, sea
nato. These were derived from expressed sequence tags (ESTs)
ism of interest. They can be used in diagnostics,
e mapping, identification of mutations, to assess
and for nutritional purposes. The present sequence is a
invention

A;

2.8%; Score 8; DB 4; Length 430;

100.0%; Pred. No. 1.6e+02;

nservative 0; Mismatches 0; Indels 0; Gaps 0;

ALAL 61

|||

ALAL 9

hard; protein; 431 AA.

first entry)

sequence Seq ID508 related to grain filling.

ology; carbohydrate synthesis; carbohydrate metabolism;
egradation; carbohydrate; plant grain; grain filling; corn;
; canola; cotton; peanut; sorghum; tobacco; sugarbeet;
rotein; oil; starch; fibre; moisture content; cereal grain;
t.

2.

002WO-IB002450.

001US-0300112P.

001US-032527P.

001US-0342327P.

TA PARTICIPATIONS AG.

W, Briggs S, Cooper B, Goff SA, Moughamer T;
Katagiri F, Krops J, Provart N, Ricke D;

XX

DR WPI; 2003-229341/22.

DR N-PSDB; ADC08202.

XX

New plant genes encoding polypeptides having an activity involve
associated with the synthesis, metabolism or degradation of carb
in the plant grain useful in generating plants having improved
nutritional properties.

XX

Claim 34; SEQ ID NO 508; 130pp; English.

XX

This invention, in the area of plant biotechnology, relates to ne
polynucleotides comprising a nucleotide sequence encoding a prote
is involved in or associated with the synthesis, metabolism or
degradation of carbohydrates in the plant grain and the expres
which is up-regulated during grain filling. The plant is selected
corn, tomato, banana, canola, cotton, peanut, sorghum, tobacco,
sugarbeet, wheat, and rice. The invention may be useful for the
improvement of protein, oil, starch, fibre and moisture content c
cereal grains. In addition, carbohydrate levels may be modified t
desirable level using the present invention. The present sequence
amino acid sequence of a rice protein of the invention. Note: The
sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly fro
at ftp.wipo.int/pub/publishedpct_sequences.

XX

Sequence 431 AA;

SQ

Query Match

Best Local Similarity 2.8%; Score 8; DB 7; Length 431;

Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY

42 QRRRRGG 49

Db

372 QRRRRGG 379

RESULT 62

ADC64562

ID ADC64562 standard; protein; 431 AA.

XX

AC ADC64562;

XX

01-JAN-2004 (first entry)

XX

Synechococcus sp. Synwh0268 protein.

XX

Plant growth; commercial yield; plant breeding; fruit yield;

KW

flowering rate; Synwh0268.

XX

Synechococcus sp.; WH 8102.

OS

US2003192076-A1.

XX

09-OCT-2003.

XX

10-APR-2003; 2003US-00410432.

XX

26-MAR-2002; 2002WO-IL000250.

XX

(YISS) YISSUM RES DEV CO HEBREW UNIV JERUSALEM.

XX

Kaplan A, Lieman-Hurwitz J, Schatz D, Mittler R, Rachmilevitch

XX

WPI; 2003-831832/77.

XX

Obtaining plants having enhanced growth and/or fruit yield and/or

PT

flowering rate, specifically C3 plants grown under limiting condit

XX

useful in plant molecular biology and commercial plant breeding.

XX

Claim 2; Fig 11; 47pp; English.

XX

The present invention relates to a method of obtaining plants with

with and/or commercial yield under growth limiting conditions. comprises obtaining a population of plants transformed to a polypeptide having at least 60% sequence identity to any of 8 sequences, growing the plants and selecting plants having the polypeptide. The methods and compositions of the present invention are useful in commercial plant breeding, particularly for plants having enhanced growth and/or fruit yield and/or ornamental value. The present sequence represents *Synechococcus* sp. protein.

AA;

2.8%; Score 8; DB 7; Length 431;
 Identity 100.0%; Pred. No. 1.6e+02;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LG LLL 74

|||||

LG LLL 371

standard; protein; 454 AA.

(first entry)

elanogaster polypeptide SEQ ID NO 516.

developmental biology; cell signalling; insecticide;
 al.

elanogaster.

A2.

2001WO-US009231.

2000US-0191637P.

2000US-00614150.

DRP NY.

Adams M, Li PWD, Myers EW;

5860/75.

2011.

nucleic acid detection reagent for detecting 1000 or more
 Drosophila and for elucidating cell signaling and cell-cell

SEQ ID NO 516; 21pp + Sequence Listing; English.

relates to an isolated nucleic acid detection reagent
 detecting 1000 or more genes from *Drosophila*. The invention is
 experimental biology and in elucidating cell signalling and
 interactions in higher eukaryotes for the development of
 therapeutics and pharmaceutical drugs. The invention
 genomic DNA sequences (ABL16176-ABL30511), expressed DNA
 H01840-ABL16175) and the encoded proteins (ABBS57737-
 ie sequence data for this patent did not form part of the
 publication, but was obtained in electronic format directly
 ftp.wipo.int/pub/published_pct_sequences

AA;

2.8%; Score 8; DB 4; Length 454;
 Identity 100.0%; Pred. No. 1.7e+02;

Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 56 LVPLALGL 63

|||||

Db 170 LVPLALGL 177

RESULT 64

ADA54710

ID ADA54710 standard; protein; 472 AA.

XX

AC ADA54710;

XX

DT 20-NOV-2003 (first entry)

XX

DE Human protein, SEQ ID 2278.

XX

KW Cytostatic; Anti-inflammatory; Osteopathic; Neuroprotective; Noc
 Gene Therapy; human; secretory protein; membrane proteins; cancer
 inflammatory disease; osteoporosis; neurological disease.

XX

OS Homo sapiens.

XX

PN EP1293569-A2.

XX

PD 19-MAR-2003.

XX

PF 21-MAR-2002; 2002EP-00006586.

XX

PR 14-SEP-2001; 2001JP-00328381.

XX

PR 24-JAN-2002; 2002US-0350435P.

XX

(HELI-) HELIX RES INST.

(REAS-) RES ASSOC BIOTECHNOLOGY.

XX

Isogai T, Sugiyama T, Otsuki T, Wakamatsu A, Sato H, Ishii
 Yamamoto J, Isono Y, Hio Y, Otsuka K, Nagai K, Irie R, Tam
 Seki N, Yoshikawa T, Otsuka M, Nagahara K, Masuho Y;

XX

WPI: 2003-395539/38.

DR N-PSDB; ADA53071.

XX

New polynucleotides encoding full-length polypeptides, e.g. secretory
 and/or membrane proteins, useful for developing medicines for di
 which the gene is involved, or as target molecules for gene ther.

XX

Claim 14; SEQ ID NO 2278; 205pp; English.

XX

The present invention relates to novel human secretory or membra
 proteins (ADA54072-ADA55710) and their coding sequences (ADA5243
 ADA54071). The coding sequences are useful in the gene therapy o
 diseases caused by abnormalities of the proteins, e.g. cancer,
 inflammatory diseases, osteoporosis or neurological disease.

XX

SQ Sequence 472 AA;

Query Match

Best Local Similarity 2.8%; Score 8; DB 6; Length 472;

Matches 8; Conservative 0; Mismatches 0; Indels 0; (

QY 60 ALGLGLAL 67

|||||

Db 440 ALGLGLAL 447

RESULT 65

ABG20260

ID ABG20260 standard; protein; 586 AA.

XX

AC ABG20260;

XX

DT 18-FEB-2002 (first entry)

XX

06:25:19 2004

us-09-245-198a-4.oligo.rag

diagnostic protein #20251.

some mapping; gene therapy; forensic;
nt; medical imaging; diagnostic; genetic disorder.

2.

2001WO-US008631.

2000US-00540217.

2000US-00649167.

INC.

Liu C, Tang YT;

362/73.

447.

polynucleotide and encoded polypeptides, useful in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits and to assess

ID NO 50619; 103pp; English.

relates to isolated polynucleotide (I) and polypeptide (II)
is useful as hybridisation probes, polymerase chain
primers, oligomers, and for chromosome and gene mapping,
inant production of (II). The polynucleotides are also used
as expressed sequence tags for identifying expressed
useful in gene therapy techniques to restore normal
II) or to treat disease states involving (II). (II) is
nerating antibodies against it, detecting or quantitating a
tissue, as molecular weight markers and as a food
II) and its binding partners are useful in medical imaging
essing (II). (I) and (II) are useful for treating disorders
rant protein expression or biological activity. The
ad polynucleotide sequences have applications in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits to assess biodiversity
e other types of data and products dependent on DNA and
quences. ABG0010-ABG30377 represent novel human diagnostic
ences of the invention. Note: The sequence data for this
appear in the printed specification, but was obtained in
mat directly from WIPO at
pub/published_pct_sequences

AA;

2.8%; Score 8; DB 4; Length 586;
arity 100.0%; Pred. No. 2.2e+02;
nservative 0; Mismatches 0; Indels 0; Gaps 0;

ALA 68

|||||

ALA 21

iard; protein; 586 AA.

(first entry)

P36269, SEQ ID NO 11960.

KW

Human; pain; neuronal tissue; gene therapy;

KW

spinal segmental nerve injury; chronic constriction injury; CCI;

KW

spared nerve injury; SNI; Chung.

XX

Homo sapiens.

XX

WO2003016475-A2.

XX

27-FEB-2003.

XX

14-AUG-2002; 2002WO-US025765.

XX

14-AUG-2001; 2001US-0312147P.

PR

01-NOV-2001; 2001US-0346382P.

PR

26-NOV-2001; 2001US-0333347P.

XX

(GEHO) GEN HOSPITAL CORP.

PA

(FARB) BAYER AG.

XX

Woolf C, D'urso D, Befort K, Costigan M;

XX

WPI; 2003-268312/26.

DR

GENBANK; P36269.

XX

New composition comprising two or more isolated polypeptides, use
preparing a medicament for treating pain in an animal.

PT

XX

Claim 1; Page; 1017pp; English.

PS

XX

The invention discloses a composition comprising two or more isol
or human polynucleotides or a polynucleotide which represents a f
derivative or allelic variation of the nucleic acid sequence. Als
claimed are a vector comprising the novel polynucleotide, a host
comprising the vector, a method for identifying a nucleotide seq
which is differentially regulated in an animal subjected to pain
kit to perform the method, an array, a method for identifying an
that increases or decreases the expression of the polynucleotide
that is differentially expressed in neuronal tissue of a first ar
subjected to pain, a method for identifying a compound which regu
the expression of a polynucleotide sequence which is differential
expressed in an animal subjected to pain, a method for identifyin
compound that regulates the activity of one or more of the
polynucleotides, a method for producing a pharmaceutical composi
method for identifying a compound or small molecule that regulate
activity in an animal of one or more of the polypeptides given in
specification, a method for identifying a compound useful in trea
pain and a pharmaceutical composition comprising the one or more
polypeptides or their antibodies. The polynucleotide or the compo
modulates its activity is useful for preparing a medicament for t
pain (e.g. spinal segmental nerve injury (SNI)), chronic constri
injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. ge
therapy). The sequence presented is a human protein (shown in Tab
the specification) which is differentially expressed during pain.
The sequence data for this patent did not form part of the printe
specification, but was obtained in electronic form directly from
ftp.wipo.int/pub/published_pct_sequences.

SQ Sequence 586 AA;

Query Match

Best Local Similarity 2.8%; Score 8; DB 7; Length 586;

Matches 8; Conservative 0; Mismatches 0; Indels 0; G

QY 61 LGUGLALA 68

Db 14 LGUGLALA 21

RESULT 67

ADE62980

ID ADE62980 standard; protein; 586 AA.

XX

AC ADE62980;

(first entry)
 n P35269, SEQ ID NO 8914.
 neuronal tissue; gene therapy;
 ntal nerve injury; chronic constriction injury; CCI;
 injury; SNI; Chung.
 -A2.
 2002WO-US025765.
 2001US-0312147P.
 2001US-0346382P.
 2001US-0333347P.
 HOSPITAL CORP.
 R AG.
 urso D, Befort K, Costigan M;
 8312/26.
 269.
 ion comprising two or more isolated polypeptides, useful for
 medicament for treating pain in an animal.
 e; 1017pp; English.
 n discloses a composition comprising two or more isolated rat
 ynucleotides or a polynucleotide which represents a fragment,
 r allelic variation of the nucleic acid sequence. Also
 a vector comprising the novel polynucleotide, a host cell
 e vector, a method for identifying a nucleotide sequence
 ferentially regulated in an animal subjected to pain and a
 m the method, an array, a method for identifying an agent
 es or decreases the expression of the polynucleotide sequence
 arentially expressed in neuronal tissue of a first animal
 pain, a method for identifying a compound which regulates
 on of a polynucleotide sequence which is differentially
 an animal subjected to pain, a method for identifying a
 regulates the activity of one or more of the
 ies, a method for producing a pharmaceutical composition, a
 identifying a compound or small molecule that regulates the
 an animal of one or more of the polypeptides given in the
 a, a method for identifying a compound useful in treating
 armaceutical composition comprising the one or more
 or their antibodies. The polynucleotide or the compound that
 s activity is useful for preparing a medicament for treating
 inal segmental nerve injury (Chung), chronic constriction
 and spared nerve injury (SNI)) in an animal (e.g. gene
 sequence presented is a human protein (shown in Table 2 of
 ition) which is differentially expressed during pain. Note:
 data for this patent did not form part of the printed
 i, but was obtained in electronic form directly from WIPO at
 'pub/published_pct_sequences.
 AA;
 2.8%; Score 8; DB 7; Length 586;
 arity 100.0%; Pred. No. 2.2e+02;
 Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 ILALA 68
 ||||
 ILALA 21
 RESULT 68
 AAU32148
 ID AAU32148 standard; protein; 592 AA.
 XX
 AC AAU32148;
 XX
 DT 18-DEC-2001 (first entry)
 XX
 DE Novel human secreted protein #2639.
 XX
 KW Human; vaccination; gene therapy; nutritional supplement;
 KW stem cell proliferation; haematopoiesis; nerve tissue regenerati
 KW immune suppression; immune stimulation; anti-inflammatory; leuk
 XX
 OS Homo sapiens.
 XX
 PN WO200179449-A2.
 XX
 PD 25-OCT-2001.
 XX
 PF 16-APR-2001; 2001WO-US008656.
 XX
 PR 18-APR-2000; 2000US-00552929.
 PR 28-JAN-2001; 2001US-00770160.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Tang YT, Liu C, Drmanac RT;
 XX
 DR WPI; 2001-611725/70.
 XX
 PT Nucleic acids encoding a range of human polypeptides, useful in
 PT vaccination, testing and therapy.
 XX
 PS Claim 20; Page 562-563; 765pp; English.
 XX
 CC The invention relates to novel human secreted polypeptides. The
 CC polypeptides and antibodies to the polypeptides are useful for
 CC determining the presence of or predisposition to a disease assoc
 CC with altered levels of polypeptide. The polypeptides are also us
 CC identifying agents (agonists and antagonists) that bind to them.
 CC expressing the proteins are useful for identifying a therapeutic
 CC for use in treatment of a pathology related to aberrant expressi
 CC physiological interactions of the polypeptide. Vectors compris
 CC nucleic acids encoding the polypeptides and cells genetically en
 CC to express them are also useful for producing the proteins. The
 CC are useful in genetic vaccination, testing and therapy, and can
 CC as nutritional supplements. They may be used to increase stem ce
 CC proliferation; to regulate haematopoiesis; and in bone, cartilag
 CC and/or nerve tissue growth or regeneration; immune suppression a
 CC stimulation; as anti-inflammatory agents; and in treatment of le
 CC AAU29510-AAU3304 represent the amino acid sequences of novel hu
 CC secreted proteins of the invention
 XX
 SQ Sequence 592 AA;
 Query Match 2.8%; Score 8; DB 4; Length 592;
 Best Local Similarity 100.0%; Pred. No. 2.2e+02;
 Matches 8; Conservative 0; Mismatches 0; Indels 0;
 QY 61 LGLGLALA 68
 |||||
 DB 44 LGLGLALA 51
 RESULT 69
 ABG03722
 ID ABG03722 standard; protein; 603 AA.
 XX
 AC ABG03722;
 XX
 DT 13-FEB-2002 (first entry)
 XX

agnostic protein #3713.
some mapping; gene mapping; gene therapy; forensic;
it; medical imaging; diagnostic; genetic disorder.
.
.
2001WO-US008631.
2000US-00540217.
2000US-00649167.
INC.
Jiu C, Tang YT;
62/73.
109.
polynucleotide and encoded polypeptides, useful in
orensics, gene mapping, identification of mutations
r genetic disorders or other traits and to assess
ID NO 34081; 103pp; English.
relates to isolated polynucleotide (I) and polypeptide (II)
is useful as hybridisation probes, polymerase chain
primers, oligomers, and for chromosome and gene mapping,
nant production of (II). The polynucleotides are also used
as expressed sequence tags for identifying expressed
useful in gene therapy techniques to restore normal
I) or to treat disease states involving (II). (II) is
erating antibodies against it, detecting or quantitating a
tissue, as molecular weight markers and as a food
I) and its binding partners are useful in medical imaging
ssing (II). (I) and (II) are useful for treating disorders
rant protein expression or biological activity. The
d polynucleotide sequences have applications in
orensics, gene mapping, identification of mutations
r genetic disorders or other traits to assess biodiversity
other types of data and products dependent on DNA and
ences. ABG00010-ABG30377 represent novel human diagnostic
ences of the invention. Note: The sequence data for this
appear in the printed specification, but was obtained in
mat directly from WIPO at
ub/published_pct_sequences
A;
2.8%; Score 8; DB 4; Length 603;
rity 100.0%; Pred. No. 2.3e+02;
nservative 0; Mismatches 0; Indels 0; Gaps 0;
ALA 68
|||
ALA 447
ard; protein; 617 AA.
first entry)
modification and maintenance molecule (PMMW)-44.

protein modification and maintenance molecule; PMMW;
protein modification; protein maintenance; protein function;
protein conformation; protein stabilisation; protein degradation;
phosphatase; protease; protease inhibitor; isomerase; transferase
molecular chaperone; anti-HIV; anti-allergic; anti-inflammatory;
antianemic; antiparkinsonian; nootropic; anticonvulsant;
antiarteriosclerotic; antiasthmatic; immunosuppressive; antithyro
cytostatic; hepatotropic; dermatological; antidiabetic; nephropro
antigout; thyromimetic; neuroprotective; osteopathic; antiarthritis
antiparasitic; antihelminthic; antipeptidic; uropathic; ophthalmic
antineumatic; haemostatic; antibacterial; virucide; protozoacide
fungicide; gene therapy; cell proliferative disorder; arterioscle
hepatitis; polycythaemia vera; psoriasis; primary thrombocytopen
cancer; developmental disorder; anaemia; mental retardation;
neurological disorder; Alzheimer's disease; Parkinson's disease;
epilepsy; autoimmune disorder; inflammatory disorder; AIDS; aller
g asthma; autoimmune thyroiditis; Crohn's disease; diabetes mellitu
glomerulonephritis; Goodpasture's syndrome; multiple sclerosis;
arthritis; osteoporosis; pancreatitis; Sjogren's syndrome;
microbial infection; human.
Homo sapiens.
WO2003063688-A2.
07-AUG-2003.
23-JAN-2003; 2003WO-US002500.
25-JAN-2002; 2002US-0351928P.
25-FEB-2002; 2002US-0359903P.
21-MAR-2002; 2002US-0366837P.
(INCY-) INCYTE GENOMICS INC.
Hafalia AJA, Li JX, Gorvad AE, Chawla NK, Sprague WW, Lee SY,
Chang H, Elliott VS, Ramkumar J, Khare R, Emerling BM, Kable
Tang YT, Yue H, Gietzen KJ, Lee S, Swarnakar A, Baughn MR;
Wilson AD, Jin P, Chien D, Hawkins PR, Jiang X, Jackson AA;
Bhatia U, Burrill JD, Blake JJ, Ho A, Zheng W, Ison CH, Marc
Tran UK, Lal PG, Warren BA, Xu Y, Honchell CD, Becha SD;
Lehr-Mason PM;
WPI: 2003-636761/60.
N-PSDB; ADE79064.
New human protein modification and maintenance molecules and
polynucleotides, useful for diagnosing, treating or preventing aut
or inflammatory disorders (e.g. AIDS, allergy or anemia), multiple
sclerosis or cancer.
Claim 1; SEQ ID NO 44; 405pp; English.
This invention relates to novel isolated human proteins, which are
protein modification and maintenance molecules (PMMW). The cellula
processes regulating modification and maintenance of protein molec
coordinate their function, conformation, stabilisation and degrada
Each of these processes is mediated by key enzymes or proteins suc
kinases, phosphatases, proteases, protease inhibitors, isomerases,
transferases and molecular chaperones. Compounds which modulate th
proteins of the invention may have anti-HIV, anti-allergic,
anti-inflammatory, antianemic, antiparkinsonian, nootropic,
anticonvulsant, antiarteriosclerotic, antiasthmatic, immunosuppres
antithyroid, cytostatic, hepatotropic, dermatological, antidiabeti
nephrotropic, antigout, thyromimetic, neuroprotective, osteopathi
antiarthritis, antiparasitic, antihelminthic, antipeptidic, uropat
ophthalmological, antirheumatic, haemostatic, antibacterial, viruc
protozoacide or fungicide activities. The DNA sequence which encod
proteins of the invention may be useful for gene therapy. The huma
protein modification and maintenance molecules (PMMWs), the DNA se
which encode them and their modulating compounds are useful for
diagnosing, treating or preventing disorders associated with aberr
expression of PMMW, particularly cell proliferative disorders (for

riosclerosis, hepatitis, polycythemia vera, psoriasis, monocytopenia or cancer), developmental disorders (for mild or mental retardation), neurological disorders (for Alzheimer's disease, Parkinson's disease or epilepsy), inflammatory disorders (for example AIDS, allergies, asthma, myeloiditis, Crohn's disease, diabetes mellitus, arthritis, Goodpasture's syndrome, multiple sclerosis, steoporosis, pancreatitis, Sjogren's syndrome) or microbial infection. The present sequence is the amino acid sequence of a human invention.

AA;

2.8%; Score 8; DB 7; Length 617;
Identity 100.0%; Pred. No. 2.3e+02;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GLALA 68

|||||

GLALA 21

standard; protein; 633 AA.

(first entry)

diagnostic protein #20252.

osome mapping; gene mapping; gene therapy; forensic;
ant; medical imaging; diagnostic; genetic disorder.

A2.

2001WO-US008631.

2000US-00540217.

2000US-00649167.

) INC.

Liu C, Tang YT;

362/73.

448.

polynucleotide and encoded polypeptides, useful in
forensics, gene mapping, identification of mutations
or genetic disorders or other traits and to assess

ID NO 50620; 103pp; English.

relates to isolated polynucleotide (I) and polypeptide (II)
is useful as hybridisation probes, polymerase chain
primers, oligomers, and for chromosome and gene mapping,
inant production of (II). The polynucleotides are also used
as expressed sequence tags for identifying expressed
useful in gene therapy techniques to restore normal
II) or to treat disease states involving (II). (II) is
nerating antibodies against it, detecting or quantitating a
n tissue, as molecular weight markers and as a food
II) and its binding partners are useful in medical imaging
essing (II). (I) and (II) are useful for treating disorders
rant protein expression or biological activity. The
nd polynucleotide sequences have applications in

diagnostics, forensics, gene mapping, identification of mutation
responsible for genetic disorders or other traits to assess bio
and to produce other types of data and products dependent on DNA
amino acid sequences. ABC0010-ABG30377 represent novel human di
amino acid sequences. Note: The sequence data f
patent did not appear in the printed specification, but was obta
CC electronic format directly from WIPO at
ftp.wipo.int/pub/published_pct_sequences

XX Sequence 633 AA;

Query Match

Best Local Similarity 2.8%; Score 8; DB 4; Length 633;

Matches 8; Conservative 0; Mismatches 0; Indels 0;

QY 61 LGLGLALA 68

|||||

Db 44 LGLGLALA 51

RESULT 72

ABG28291

ID ABG28291 standard; protein; 799 AA.

XX ABG28291;

XX 18-FEB-2002 (first entry)

XX Novel human diagnostic protein #28282.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder.
OS Homo sapiens.

XX WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US008631.

XX 31-MAR-2000; 2000US-00540217.

XX 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX N-PSDB; AAS92478.

XX New isolated polynucleotide and encoded polypeptides, useful in

diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity.

XX Claim 20; SEQ ID NO 58650; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypept

sequences. (I) is useful as hybridisation probes, polymerase chain
reaction (PCR) primers, oligomers, and for chromosome and gene ma
and in recombinant production of (II). The polynucleotides are al
in diagnostics as expressed sequence tags for identifying express
genes. (I) is useful in gene therapy techniques to restore normal
activity of (II) or to treat disease states involving (II). (II)
useful for generating antibodies against it, detecting or quantit
polypeptide in tissue, as molecular weight markers and as a food
supplement. (II) and its binding partners are useful in medical i
of sites expressing (II). (I) and (II) are useful for treating di
involving aberrant protein expression or biological activity. The
polypeptide and polynucleotide sequences have applications in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits to assess biodi

Other types of data and products dependent on DNA and sequences. ABG00010-ABG30377 represent novel human diagnostic sequences of the invention. Note: The sequence data for this appear in the printed specification, but was obtained in format directly from WIPO at pub/published_pct_sequences

AA;
arity 2.8%; Score 8; DB 4; Length 799;
conservative 0; Mismatches 0; Indels 0; Gaps 0;
3LAL 67
3LAL 294

ard; protein; 842 AA.

(first entry)

(useful for identifying genetic disorders) #630.

vel protein; tissue marker; molecular weight marker;
xer; genetic disorder.

2.

002WO-US039555.

001US-0339739P.
001US-0339453P.
002US-0365091P.
002US-0365384P.
002US-0372381P.
002US-0372615P.
002US-00128558.
002US-0376045P.

INC.

di V, Goodrich RW, Ren F, Zhang J, Zhao QA, Wang J;
AJ, Wehrman T, Weng G, Zhou P, Drmanac RT, Wang Z;
Chen R, Xu C, Boyle BJ;

35/53.

64.

tides, useful for expressing recombinant proteins for
acterization or therapeutic use, or as markers for tissues
corresponding protein is preferentially expressed.

ID NO 1541; 1177pp; English.

comprises the amino acid and coding sequences of novel
DNA and protein sequences of the invention are useful as:
issues in which the corresponding protein is preferentially
molecular weight markers on gels; as chromosome markers or
ify chromosomes or to map related gene positions; and to
ndogenous DNA sequences in patients to identify potential
ers. The present amino acid sequence represents a protein
on.

A;

Query Match 2.8%; Score 8; DB 7; Length 842;
Best Local Similarity 100.0%; Pred. No. 3.1e+02;
Matches 8; Conservative 0; Mismatches 0; Indels 0;

Oy 54 ALLVPLAL 61
Db 441 ALLVPLAL 448

RESULT 74

ABU16705
ID ABU16705 standard; protein; 1032 AA.

XX AC ABU16705;

XX DT 19-JUN-2003 (first entry)

XX DE Protein encoded by Prokaryotic essential gene #2232.

XX KW Antisense; prokaryotic essential gene; cell proliferation; drug d

XX OS Acinetobacter baumannii.

XX PN WO200277183-A2.

XX PD 03-OCT-2002.

XX PF 21-MAR-2002; 2002WO-US009107.

XX PR 21-MAR-2001; 2001US-00815242.

XX PR 06-SEP-2001; 2001US-00948993.

XX PR 25-OCT-2001; 2001US-0342923P.

XX PR 08-FEB-2002; 2002US-00072851.

XX PR 06-MAR-2002; 2002US-0362699P.

XX PA (ELIT-) ELITRA PHARM INC.

XX PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind

XX PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

XX XX WPI; 2003-029926/02.

XX DR N-PSDB; ACA20575.

XX DR New antisense nucleic acids, useful for identifying proteins or s

XX PT for homologous nucleic acids required for cellular proliferation

XX PT isolate candidate molecules for rational drug discovery programs.

XX PS Claim 25; SEQ ID NO 44629; 1766pp; English.

XX CC The invention relates to an isolated nucleic acid comprising any

XX CC of the 6213 antisense sequences given in the specification where exp

XX CC of the nucleic acid inhibits proliferation of a cell. Also include

XX CC (1) a vector comprising a promoter operably linked to the nucleic

XX CC encoding a polypeptide whose expression is inhibited by the antise

XX CC nucleic acid; (2) a host cell containing the vector; (3) an isolat

XX CC polypeptide or its fragment whose expression is inhibited by the

XX CC antisense nucleic acid; (4) an antibody capable of specifically bi

XX CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cell

XX CC proliferation or the activity of a gene in an operon required for

XX CC proliferation; (7) identifying a compound that influences the acti

proliferation to isolate candidate molecules for rational
 ry programs, or for screening homologous nucleic acids
 proliferation in cells other than *S. aureus*, *S. typhimurium*,
 e or *P. aeruginosa*. The present sequence is encoded by one of
 rokaryotic essential genes. Note: The sequence data for this
 ot form part of the printed specification, but was obtained
 c format directly from WIPO at
 /pub/published_pct_sequences

2 AA;

2.8%; Score 8; DB 6; Length 1032;
 larity 100.0%; Pred. No. 3.8e+02;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

ALGLG 64

|||||

ALGLG 981

ndard; protein; 1033 AA.

(first entry)

r baumannii protein #1012.

r baumannii; bacterial disease; antibacterial; vaccine;
 crol agent.

r baumannii.

99US-00328352.

98US-0088701P.

4E THERAPEUTICS CORP.

ish D;

5092/54.

1725.

acter baumannii proteins and nucleic acids, useful as reagents
 ing a bacterial disease, as components of antibacterial
 targets for antibacterial drugs, or as biocontrol agents for

ID NO 5138; 328pp; English.

i relates to isolated Acinetobacter baumannii nucleic acids.
 uni nucleic acids and polypeptides are useful as reagents
 ing a bacterial disease, as components of antibacterial
 targets for antibacterial drugs, to detect the presence of
 and other Acinetobacter species in a sample, in screening
 the ability to interfere with the A. baumannii life cycle
 : A. baumannii infection, and as biocontrol agents for
 present sequence represents the amino acid sequence of an A.
 otein.

AA;

2.8%; Score 8; DB 6; Length 1033;

arity 100.0%; Pred. No. 3.8e+02;

Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 57 VPLALGLG 64
 |||||

Db 975 VPLALGLG 982

Search completed: April 7, 2004, 17:57:27
 Job time : 65 secs